## AvMed

## 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 1-877-535-1391</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>.

For Medicare Members: Medicare Coverage for outpatient (Part B) drugs is outlined in the Medicare Benefit Policy Manual (Pub. 100-2), Chapter 15, §50 Drugs and Biologicals. In addition, National Coverage Determination (NCD) and Local Coverage Determinations (LCDs) may exist and compliance with these policies is required where applicable. They can be found at: <a href="https://www.cms.gov/medicare-coverage-database/overview-and-quick-search.aspx">https://www.cms.gov/medicare-coverage-database/overview-and-quick-search.aspx</a>. Additional indications may be covered at the discretion of the health plan.

Drug Requested: Sunlenca® (lenacapavir) (J1961) (Medical)

MEMBER & PRESCRIBER INFO	RMATION: Authorization may be delayed if incomplete.
Member Name:	
Member AvMed #:	
Prescriber Name:	
Prescriber Signature:	Date:
Office Contact Name:	
Phone Number:	Fax Number:
NPI #:	
DRUG INFORMATION: Authorizatio	
Drug Name/Form/Strength:	
Dosing Schedule:	Length of Therapy:
Diagnosis:	ICD Code, if applicable:
Weight (if applicable):	Date weight obtained:
	e timeframe does not jeopardize the life or health of the member in function and would not subject the member to severe pain.
Maintenance Dose: 927 mg by subcutaned from the date of the last injection +/- 2 weeks	ous injection (2 x 1.5 mL injections) every 6 months (26 weeks)
<b>Quantity Limit:</b> 3 mL per 184 days	
	all that apply. All criteria must be met for approval. To , including lab results, diagnostics, and/or chart notes, must be
☐ Member is ≥ 12 years of age and weight	hing $\geq$ 35 kg, or an adult aged $\geq$ 18 years
☐ Prescribed by, or in consultation with,	an infectious disease specialist or specialist in HIV treatment

(Continued on next page)

Member has been identified to have multidrug resistant HIV-1 infection with documented resistance
to at least $\underline{\text{TWO}}$ (2) antiretroviral medications from $\geq 3$ of the 4 main antiretroviral drug classes below
(must submit genotype/phenotype resistance testing results):
□ Nucleoside Reverse Transcriptase Inhibitors/Non-nucleoside Reverse Transcriptase Inhibitors
□ Protease Inhibitors
☐ Entry Inhibitors (including CCR5 antagonists)
☐ Integrase Inhibitor
Member is experiencing current virologic failure defined as having a viral load greater than 400 copies/mL before treatment initiation
Member's current viral load has been submitted with request
• Current Viral Load:copies/mL (must submit most recent labwork indicating viral load prior to initiating therapy, within 4-8 weeks)
Provider confirms requested medication will be used in conjunction with an optimized background regimen for antiretroviral therapy
Provider confirms requested medication will be initiated using <b>ONE</b> of the following dosing regimens:

□ Initiation Option 1			
Day 1	927 mg by subcutaneous injection (2 x 1.5 mL injections) <u>AND</u> 600 mg orally (2 x 300 mg tablets)		
Day 2	600 mg orally (2 x 300 mg tablets)		
□ Initiation Option 2			
Day 1	600 mg orally (2 x 300 mg tablets)		
Day 2	600 mg orally (2 x 300 mg tablets)		
Day 8	300 mg orally (1 x 300 mg tablet)		
Day 15	927 mg by subcutaneous injection (2 x 1.5 mL injections)		

Medication being provided by (check applicable box(es) below):				
□ Physician's office	OR	□ Specialty Pharmacy – PropriumRy		

\*\*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. \*