

# SENTARA HEALTH PLANS

## PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST\*

**Directions:** 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; **1-800-750-9692**. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. **If the information provided is not complete, correct, or legible, the authorization process can be delayed.**

**Drug Requested:** Sunlenca<sup>®</sup> (lenacapavir) (Pharmacy)

**MEMBER & PRESCRIBER INFORMATION:** Authorization may be delayed if incomplete.

Member Name: \_\_\_\_\_

Member Sentara #: \_\_\_\_\_ Date of Birth: \_\_\_\_\_

Prescriber Name: \_\_\_\_\_

Prescriber Signature: \_\_\_\_\_ Date: \_\_\_\_\_

Office Contact Name: \_\_\_\_\_

Phone Number: \_\_\_\_\_ Fax Number: \_\_\_\_\_

DEA OR NPI #: \_\_\_\_\_

**DRUG INFORMATION:** Authorization may be delayed if incomplete.

Drug Form/Strength: \_\_\_\_\_

Dosing Schedule: \_\_\_\_\_ Length of Therapy: \_\_\_\_\_

Diagnosis: \_\_\_\_\_ ICD Code: \_\_\_\_\_

Weight: \_\_\_\_\_ Date: \_\_\_\_\_

**Maintenance Dose:** 927 mg by subcutaneous injection (2 x 1.5 mL injections) every 6 months (26 weeks) from the date of the last injection +/- 2 weeks

**Quantity Limit:** 3 mL per 184 days

**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.

- Member is  $\geq 12$  years of age and weighing  $\geq 35$  kg, or an adult aged  $\geq 18$  years
- Prescribed by, or in consultation with, an infectious disease specialist or specialist in HIV treatment
- Member has been identified to have multidrug resistant HIV-1 infection with documented resistance to at least **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

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- 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 **ONE** of the following dosing regimens:

<input type="checkbox"/> <b>Initiation Option 1</b>	
Day 1	927 mg by subcutaneous injection (2 x 1.5 mL injections) <b><u>AND</u></b> 600 mg orally (2 x 300 mg tablets)
Day 2	600 mg orally (2 x 300 mg tablets)
<input type="checkbox"/> <b>Initiation Option 2</b>	
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 Specialty Pharmacy - PropriumRx**

***\*\*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.\****