



FLORIDA MEDICAID PRIOR AUTHORIZATION

Fuzeon®

(Maximum Length of Approval is 6 Months)

Note: Form must be completed in full. An incomplete form may be returned.

Recipient's Medicaid ID#

Grid for Recipient's Medicaid ID#

Date of Birth (MM/DD/YYYY)

Grid for Date of Birth (MM/DD/YYYY)

Recipient's Full Name

Grid for Recipient's Full Name

Prescriber's Full Name

Grid for Prescriber's Full Name

Prescriber's NPI

Grid for Prescriber's NPI

Prescriber Phone Number

Grid for Prescriber Phone Number

Prescriber Fax Number

Grid for Prescriber Fax Number

Pharmacy Name

Grid for Pharmacy Name

Pharmacy Medicaid Provider #

Grid for Pharmacy Medicaid Provider #

Pharmacy Phone Number

Grid for Pharmacy Phone Number

Pharmacy Fax Number

Grid for Pharmacy Fax Number

Drug: _____ Quantity: _____

Length of Therapy on Prescription: _____ Dosage and Frequency of Dosing: _____

- 1. Initiation of therapy OR Continuation of therapy
2. Has the patient had a genotype/phenotype completed?
3. Does the patient have a viral load completed in the past 6 months?
4. Has the patient had a CD4 count completed in the past 6 months?
5. Has the patient been compliant with previous therapy?

Prescriber's Signature: _____ Date: _____

REQUIRED FOR REVIEW: All copies of medical records (e.g., diagnostic evaluations and recent chart notes), and the most recent copies of related labs. The provider must retain copies of all documentation for five years.

Fax this form to 1-866-940-7328

Pharmacy PA Call Center:

1-855-258-1593

02.15.2024

Confidentiality Notice: The documents accompanying this transmission contain confidential health information that is legally privileged. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution, or action taken in reliance on the contents of these documents is strictly prohibited.

Use with PA Form

Question 1 and 2 For initiation of therapy, genotype, and phenotype results should be dated within the past 12 months.

Note: Genotyping and phenotyping cannot be effectively done if the viral load is less than 1000 copies/mL. Therefore, genotyping and phenotyping is not required for those recipients currently on Fuzeon therapy.

Question 3 Only acceptable response for approval is “Yes.”

Question 4 Only acceptable response for approval is “Yes.”

Question 5 New therapy requires verification of:

- 1) Ongoing therapy with other HIV medications
- 2) Compliance on previous therapies
- 3) Labs that demonstrate CD4 counts and antigen levels consistent with medication failure.

Continuation of therapy requires verification of compliance with other medications. If Fuzeon is working, then CD4 counts should be good and viral antigen levels should be undetectable.

Approved Indications

Fuzeon, in combination with other antiretroviral agents, is indicated for the treatment of HIV-1 infection in treatment-experienced patients with evidence of HIV-1 replication despite ongoing antiretroviral therapy.

Approval Period

Maximum of six months.