

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.

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Prime Therapeutics State Government Solutions LLC Prior Authorization P. O. Box 7082 Tallahassee, FL 32314-7082 Phone: 877-553-7481 Fax: 877-614-1078 **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. If you have received this information in error, please notify the sender (via return fax) immediately and arrange for the return or destruction of these documents. Distribution, reproduction or any other use of this transmission by any party other than the intended recipient is strictly prohibited.



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