

Prior Authorization Request Form for Intron A/Infergen**Member Information**

Patient Name _____
 Cardholder ID _____
 Date of Birth _____
 Address _____
 City, State Zip _____
 Phone Number _____

Provider Information

Provider Name _____
 DEA Number _____
 Address _____
 City, State and Zip _____
 Phone Number _____
 FAX Number _____

Pharmacy Information

Pharmacy Name _____ Address _____ Phone _____

Drug Name and Strength _____**Criteria for Approval:**

- Does the patient have a diagnosis of: *Check all that apply:*

<input type="checkbox"/> AIDS-related Kaposi's Sarcoma	<input type="checkbox"/> Chronic Hepatitis B	<input type="checkbox"/> Chronic Hepatitis C
<input type="checkbox"/> Condyloma acuminatum (genital warts)	<input type="checkbox"/> Follicular Non-Hodgkin's Lymphoma	
<input type="checkbox"/> Hairy Cell Leukemia	<input type="checkbox"/> Malignant Melanoma	
- Has the patient received interferon therapy within the previous year? Yes No
- Did the patient respond to the previous interferon therapy? Yes No
- Did the patient have detectable serum levels of hepatic C virus (HCV) RNA after or at the end of initial treatment with an interferon? Yes No
- Does the patient have: *Check all that apply:*

<input type="checkbox"/> Detectable levels of hepatitis C virus (HBV) RNA in the serum
<input type="checkbox"/> Serum markers of hepatitis B virus (HBV) replication (HBeAg and HBV DNA)
<input type="checkbox"/> Demonstrated chronic hepatitis C on liver biopsy
<input type="checkbox"/> Persistently elevated serum alanine aminotransferase (ALT) levels > 2 times the upper limit
<input type="checkbox"/> A viral load of at least > 2-log decrease
- Is the patient Genotype-1? Yes No
- Will Intron A be used in conjunction with anthracycline containing chemotherapy? Yes No
- Will the patient be monitored for depression during therapy? Yes No
- Will the hemoglobin levels be monitored during therapy? Yes No
- Is the patient being treated for relapse after monotherapy with an interferon product? Yes No
- If the patient has relapsed or did not respond to previous interferon therapy, will the patient be restarted on a higher dose regimen as labeling recommends? Yes No

Provider Signature _____ Date _____

Fax completed forms to (866) 284-4509.

For Office Use Only

Date/Time Received _____
 Reference Number _____
 Approved / Denied (Circle One) by _____ Date _____
 Date/Time Returned to Provider _____

If you have any questions regarding this form, contact the Prior Authorization Department Toll Free at (866) 284-4492 or Fax Toll Free at (866) 284-4509.

FOX Rx Care Utilization Management
 3375-I Capital Circle NE
 Tallahassee, FL 32308

IMPORTANT NOTICE: This facsimile is intended to be delivered to the named addressee and may contain material that is confidential, privileged, proprietary or exempt from disclosure and applicable law. If it is received by anyone other than the named addressee, the recipient should immediately notify the sender at the address and telephone number set forth herein and obtain instructions as to disposal of the transmitted material. In no event should such material be read or retained by other than the named addressee, except by express authority of the sender to the named addressee.

