IntronA (interferon Alfa-2b) Prior Authorization Request Form

Caterpillar Prescription Drug Benefit Phone: 877-228-7909 Fax: 800-424-7640

Instructions: Please fill out all applicable sections completely and legibly. Attach any additional documentation that is important for the review (e.g., chart notes or lab data, to support the authorization request). Information contained in this form is Protected Health Information under HIPAA.

	URGENT	
MEMBER INFORMATION		
LAST NAME:	FIRST NAME:	
PHONE NUMBER:	DATE OF BIRTH:	
STREET ADDRESS:		
CITY:	STATE: ZIP CODE:	
PATIENT INSURANCE ID NUMBER:		
MALE FEMALE HEIGHT (IN/CM): WEIGH	IT (LB/KG): ALLERGIES:	

IF YOU ARE NOT THE PATIENT OR THE PRESCRIBER, YOU WILL NEED TO SUBMIT A PHI DISCLOSURE AUTHORIZATION FORM WITH THIS REQUEST WHICH CAN BE FOUND AT THE FOLLOWING LINK: PRIMETHERAPEUTICS.COM/NOPP

PRESCRIBER INFORMATION				
LAST NAME:	FIRST NAME:			
PRESCRIBER SPECIALTY:	EMAIL ADDRESS:			
NPI NUMBER:	DEA NUMBER:			
PHONE NUMBER:	FAX NUMBER:			
STREET ADDRESS:				
CITY:	STATE: ZIP CODE:			
REQUESTOR (if different than prescriber):	OFFICE CONTACT PERSON:			

MEDICATION OR MEDICAL DISPENSING INFORMATION					
MEDICATION NAME:					
DOSE/STRENGTH:	FREQUENCY:	LENGTH OF THERAPY/REFILLS:	QUANTITY:		
NEW THERAPY		IF RENEWAL: DATE THERAPY	INITIATED:		
DURATION OF THERAPY (SPECIFIC DATES):					

Continued on next page.



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MEMBER'S LAST NAME:	MEMBER'S FIRST NAME:			
1. HAS THE PATIENT TRIED ANY OTHER	R MEDICATIONS FOR THIS CONDITION?	YES (if yes, complete below) NO		
MEDICATION/THERAPY (SPECIFY	DURATION OF THERAPY (SPECIFY	RESPONSE/REASON FOR		
DRUG NAME AND DOSAGE):	DATES):	FAILURE/ALLERGY:		
2. LIST DIAGNOSES:		ICD-10:		
AIDS-related Kaposi's sarcoma				
Chronic hepatitis B				
Chronic hepatitis C				
Condylomata acuminate				
Follicular non-Hodgkin's lymphoma				
Hairy cell leukemia				
Malignant melanoma				
Relapsed/refractory advanced cutaneous	s T-cell lymphoma*			
Renal carcinoma				
Other diagnosis:ICD-	10			
*Please provide chart documentation (i.e.,	chart notes) supporting this information.			
3. REQUIRED CLINICAL INFORMATION	: PLEASE PROVIDE ALL RELEVANT CLINIC	AL INFORMATION TO SUPPORT A		
PRIOR AUTHORIZATION.				
For chronic hepatitis B, answer the fol	lowing:			
Does the patient have compensated li	-			
Is Intron A prescribed by a gastroenterologist, infectious disease physician, hepatologist, or a transplant physician? Yes No Has the patient been serum HBsAg positive for at least 6 months with evidence of HBV replication (serum HBeAg positive)?* Yes No				
Does the patient have elevated serum ALT?* Yes No				
Does the patient have a history of hepatic encephalopathy, variceal bleeding, ascites, or other clinical signs of decompensation? Yes No 				
Is the patient's bilirubin level normal?* u Yes u No				
Are the patient's albumin levels stable and within normal limits?* \square Yes \square No				
Is the patient's prothrombin time less than 3 seconds prolonged for adults or less than or equal to 2 seconds prolonged for pediatric patients?* \Box Yes \Box No				
Is the patient's white blood count (WBC) greater than or equal to 4,000/mm ³ ?* \Box Yes \Box No				
Is the patient's platelet count greater than or equal to 100,000/mm ³ for adults or greater than or equal to 150,000/mm ³ for pediatric patients?* Yes No *Please provide lab documentation				



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For <u>chronic hepatitis C</u>, answer the following: Does the patient have compensated liver disease? □ Yes □ No

Is Intron A prescribed by a gastroenterologist, infectious disease physician, hepatologist, or a transplant physician? □ Yes □ No

Does the patient have a history of hepatic encephalopathy, variceal bleeding, ascites, or other clinical signs of decompensation?

Yes
No

Is the patient's bilirubin level less than or equal to 2 mg/dL?* \square Yes \square No

Are the patient's albumin levels stable and within normal limits?* \Box Yes \Box No

Is the patient's prothrombin time less than 3 seconds prolonged?*

Yes
No

Is the patient's white blood count (WBC) greater than or equal to 3,000/mm³?* Yes
No

Is the patient's platelet count greater than or equal to 70,000/mm³?* \Box Yes \Box No *Please provide lab documentation.

For <u>condylomata acuminate</u>, answer the following: Is Intron A being used intralesionally? □ Yes □ No

Does the condition involve external surfaces of the genital and perianal area?

Yes
No

For <u>follicular non-Hodgkin's lymphoma</u>, answer the following: Is Intron A being used in conjunction with anthracycline-containing chemotherapy?

Yes
No

For <u>malignant melanoma</u>, answer the following: Is the patient free of disease but has a high risk of systemic recurrence within 56 days of surgery? □ Yes □ No

For <u>relapsed/refractory advanced cutaneous T-cell lymphoma</u>, answer the following: Reauthorization: Has the patient been tolerant of therapy and have they had a positive continued response?
Que Yes Que No

For <u>renal cell carcinoma</u>, answer the following: Is the patient using Intron A as monotherapy? \Box Yes \Box No

Is Intron A being used in combination with bevacizumab as first line therapy for relapsed or medically unresectable stage IV disease with predominant clear cell histology?
□ Yes □ No

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?



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Please note: Not all drugs/diagnosis are covered on all plans. This request may be denied unless all required information is received.

ATTESTATION: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan, insurer, Medical Group or its designees may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.

Prescriber Signature or Electronic I.D. Verification:

Date:

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 immediately (via return FAX) and arrange for the return or destruction of these documents.

FAX THIS FORM TO: 800-424-7640

MAIL REQUESTS TO: Prime Therapeutics Management Prior Authorization Program

Attn: CP - 4201 P.O. Box 64811 St. Paul, MN 55164-0811

