

**PegIntron (peginterferon 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): \_\_\_\_\_ WEIGHT (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](http://PRIMETHERAPEUTICS.COM/NOPP)

PATIENT'S AUTHORIZED REPRESENTATIVE (IF APPLICABLE): \_\_\_\_\_

AUTHORIZED REPRESENTATIVE'S PHONE NUMBER: \_\_\_\_\_

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:
<input type="checkbox"/> NEW THERAPY	<input type="checkbox"/> RENEWAL	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: \_\_\_\_\_

<b>1. HAS THE PATIENT TRIED ANY OTHER MEDICATIONS FOR THIS CONDITION?</b> <input type="checkbox"/> YES (if yes, complete below) <input type="checkbox"/> NO		
<b>MEDICATION/THERAPY (SPECIFY DRUG NAME AND DOSAGE):</b>	<b>DURATION OF THERAPY (SPECIFY DATES):</b>	<b>RESPONSE/REASON FOR FAILURE/ALLERGY:</b>
<b>2. LIST DIAGNOSES:</b>		<b>ICD-10:</b>
<input type="checkbox"/> Chronic hepatitis B <input type="checkbox"/> Chronic hepatitis C <input type="checkbox"/> Other diagnosis: _____ ICD-10: _____		
<b>3. REQUIRED CLINICAL INFORMATION: PLEASE PROVIDE ALL RELEVANT CLINICAL INFORMATION TO SUPPORT A PRIOR AUTHORIZATION.</b>		
<b>Clinical Information:</b> Has the patient had a trial and inadequate response to a 3 month trial with Pegasys? <input type="checkbox"/> Yes <input type="checkbox"/> No  Is the prescriber a gastroenterologist, infectious disease physician, hepatologist, or a transplant physician? <input type="checkbox"/> Yes <input type="checkbox"/> No		
For <u>chronic hepatitis C</u> , also answer the following: Select if the patient has a diagnosis of chronic hepatitis C which will be treated with one of the following therapy: <input type="checkbox"/> Monotherapy (PegIntron alone) <input type="checkbox"/> Dual therapy (PegIntron and ribavirin) <input type="checkbox"/> Triple therapy  For monotherapy:* Does the patient have an intolerance or contraindication to ribavirin therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No  Does the patient have a baseline (pre-treatment) HCV-RNA assessed for the diagnosis? <input type="checkbox"/> Yes <input type="checkbox"/> No <i>*Please submit documentation supporting this information.</i>  Reauthorization: Is there at least a 2 log (100 fold) decrease in the HCV RNA level at week 12 of therapy?* <input type="checkbox"/> Yes <input type="checkbox"/> No <i>*Please submit documentation supporting this information.</i>  For dual therapy: Does the patient have compensated liver disease? <input type="checkbox"/> Yes <input type="checkbox"/> No  Document the patient's genotype:* _____  Document patient's baseline (pre-treatment) HCV-RNA level:* _____ <i>*Please submit documentation supporting this information.</i>  Reauthorization: Document the patient's genotype:* _____  Select which week of therapy the patient has completed thus far: <input type="checkbox"/> 12 weeks <input type="checkbox"/> 24 weeks		

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Select the patient's current viral load:\*

- Detectable  Undetectable  
 For patients who have completed 12 weeks of therapy, less than a 2 log reduction

\*Please submit documentation supporting this information.

**For triple therapy:**

Select if the patient has a diagnosis of chronic hepatitis C virus that will be treated with triple therapy using the following medications:

- Olysio  Ribavirin  Sovaldi

For triple therapy with Victrelis, will dual therapy with peg-interferon and ribavirin be initiated 4 weeks before Victrelis is started?  Yes  No

Document the patient's genotype: \* \_\_\_\_\_

Does the patient have compensated liver disease? \*  Yes  No

\*Please submit documentation supporting this information.

**Reauthorization:**

Select if Pegasys and ribavirin will be taken with the following antivirals:

- Olysio  Sovaldi

Select if the following applies to the patient:\*

- Treatment-naïve without cirrhosis  
 Null responder on prior treatment without cirrhosis  
 Relapser on prior treatment without cirrhosis  
 Cirrhosis  
 Partial responder on prior treatment without cirrhosis

Select which week of therapy the patient has completed thus far:

- 12 weeks  24 weeks

Select if the patient has HCV RNA levels as follows:\*

- Undetectable at week 4  
 Undetectable at week 8  
 Undetectable at weeks 4 AND 12  
 1,000 IU/mL or less at week 12 of treatment  
 Undetectable at week 24

\*Please submit documentation

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