

Please complete all required sections to allow your request to be processed.

Patients may or may not meet eligibility requirements as established by Alberta Government sponsored drug programs.

| PATIENT INFORMATION | | | | COVERAGE TYPE: |
|-------------------------|--------------------------------|---------|--|---------------------------|
| PATIENT LAST NAME | FIRST NAME | INITIAL | <input type="checkbox"/> Alberta Blue Cross <input type="checkbox"/> Alberta Human Services <input type="checkbox"/> Other | |
| BIRTH DATE (YYYY-MM-DD) | ALBERTA PERSONAL HEALTH NUMBER | | | |
| STREET ADDRESS | CITY | PROV | POSTAL CODE | ID/CLIENT/COVERAGE NUMBER |

| NOTIFICATION | PATIENT CONSENT |
|--|--|
| You may be eligible to receive Pegasys drug benefits. Information from your prescriber is required to determine eligibility. Your consent is required: (A) for your prescriber to release necessary and relevant information to Alberta Blue Cross, Alberta Health and, if requested, to Alberta Human Services; and (B) for Alberta Blue Cross to release that and related usage information to Alberta Health. | I hereby authorize: (A) my prescriber to release to Alberta Blue Cross, Alberta Health, and (if they request it) to Alberta Human Services (the aforesaid being the "designated recipients"); and (B) Alberta Blue Cross to release to Alberta Health the information on this form and information relating to my usage of and experience with the drug and treatment results, and I consent to the designated recipients collecting such information. Date _____ Patient's signature _____ |

| PRESCRIBER INFORMATION | | | |
|------------------------|---|---------|---|
| PRESCRIBER LAST NAME | FIRST NAME | INITIAL | PRESCRIBER PROFESSIONAL ASSOCIATION REGISTRATION |
| STREET ADDRESS | | | <input type="checkbox"/> CPSA <input type="checkbox"/> ACO REGISTRATION NUMBER <input type="checkbox"/> CARNA <input type="checkbox"/> ADA+C <input type="checkbox"/> ACP <input type="checkbox"/> Other |
| CITY, PROVINCE | PHONE | FAX | |
| POSTAL CODE | FAX NUMBER MUST BE PROVIDED WITH EACH REQUEST SUBMITTED | | |

| Diagnosis of chronic hepatitis C | YES | NO | Not tested |
|--|--------------------------|--------------------------|--------------------------|
| Is the patient serum HCV RNA positive (by PCR), pre-treatment..... | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> |

| Evidence of active liver disease: | YES | NO | Not tested |
|--|--------------------------|--------------------------|--------------------------|
| At least one of the following a) does the patient have elevated liver enzymes (ALT and/or AST), pre-treatment OR b) does the patient have an abnormal liver biopsy (inflammation and/or fibrosis)..... OR c) does the patient have elevated liver stiffness as demonstrated by transient elastography (fibrosis)..... | <input type="checkbox"/> | <input type="checkbox"/> | <input type="checkbox"/> |

If the patient is currently on peginterferon alfa-2a, indicate start date (YYY-MM-DD):

| INITIAL REQUEST: | EXTENSION REQUEST: |
|---|--|
| Is the patient intolerant to ribavirin? <input type="checkbox"/> YES <input type="checkbox"/> NO Is a baseline serum sample stored for future testing? <input type="checkbox"/> YES <input type="checkbox"/> NO Initial length of approval <input type="checkbox"/> Advanced fibrosis or cirrhosis (regardless of genotype).....48 weeks <input type="checkbox"/> Genotype 114 weeks <input type="checkbox"/> Genotype 2 or 314 weeks <input type="checkbox"/> Genotype 4, 5 or 614 weeks | Request for treatment extension at 14 weeks (excluding patients with advanced fibrosis and cirrhosis) Is the patient serum HCV RNA negative at 12 weeks? <input type="checkbox"/> YES → Patient may be eligible for an additional 34 weeks of coverage (total 48 wks) <input type="checkbox"/> NO → Has the patient achieved a reduction of viral load by at least 2 logs (100 fold)? <input type="checkbox"/> YES → Patient may be eligible for an additional 34 weeks of coverage (total 48 wks) <input type="checkbox"/> NO |

| PREVIOUS THERAPY: Consideration may be given to patients who have previously received therapy and who meet at least one of the following |
|---|
| <input type="checkbox"/> Advanced fibrosis or cirrhosis <input type="checkbox"/> Patient relapsed following non-pegylated interferon/ribavirin combination therapy |

| Additional information relating to request | | |
|--|------|--|
| PRESCRIBER'S SIGNATURE | DATE | Please forward this request to • Alberta Blue Cross, Clinical Drug Services 10009 108 Street NW, Edmonton, Alberta T5J 3C5 • FAX: 780-498-8384 in Edmonton • 1-877-828-4106 toll free all other areas |

ONCE YOUR REQUEST HAS SUCCESSFULLY TRANSMITTED, PLEASE DO NOT MAIL OR RE-FAX YOUR REQUEST
