



OMBITASVIR/PARITAPREVIR/RITONAVIR+
DASABUVIR
SPECIAL AUTHORIZATION REQUEST FORM

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
PATIENT LAST NAME, FIRST NAME, INITIAL, DATE OF BIRTH, ALBERTA PERSONAL HEALTH NUMBER, STREET ADDRESS, CITY, PROV, POSTAL CODE, ID/CLIENT/COVERAGE NUMBER, COVERAGE TYPE

NOTIFICATION
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.
PATIENT CONSENT
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.

PRESCRIBER INFORMATION
PRESCRIBER LAST NAME, FIRST NAME, INITIAL, PRESCRIBER PROFESSIONAL ASSOCIATION REGISTRATION, STREET ADDRESS, CITY, PROVINCE, POSTAL CODE, REGISTRATION NUMBER, PHONE, FAX, FAX NUMBER MUST BE PROVIDED WITH EACH REQUEST SUBMITTED

Please note: Where special authorization criteria have been satisfied, patients will be approved for coverage of Ombitasvir/Paritaprevir/Ritonavir+Dasabuvir (e.g. Holkira Pak) alone or in combination with Ribavirin as outlined in the special authorization criteria ("Duration of therapy reimbursed" section)

Diagnosis of chronic hepatitis C

- 1) Is the patient serum HCV RNA positive (by PCR), pre-treatment? YES NO Not Tested
2) Please indicate the patient's HCV Genotype: Genotype 1a Genotype 1b Genotype 1 unknown subtype Genotype 1 mixed infection Not Tested Other (Specify)
3) Does the patient have compensated liver disease (Child Pugh score <= 6)? YES NO
4) Does the patient have detectable HCV RNA within 6 months from request? YES test date (YYYY/MM/DD) NO
5) Does the patient have a fibrosis stage of F2, F3, or F4? YES (specify stage) NO
6) Does the patient have cirrhosis? YES NO
7) If the patient is currently on ombitasvir/paritaprevir/ritonavir+dasabuvir please indicate the start date (YYYY/MM/DD)

Previous therapy: please check ALL that apply

- Treatment naive (no prior exposure to any interferon, ribavirin or other approved or experimental HCV-specific direct-acting antiviral agent at the time of treatment initiation)
Treatment experienced (previously been treated with PegINF/RBV and did NOT receive adequate response)
-> For genotype 1a cirrhotic patients ONLY, please check one of the following
Prior relapse Partial responder Prior null response Other (please specify)
Other (please specify)

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

The information on this form is being collected and pursuant to sections 20, 21 and 22 of the Health Information Act, and sections 33 and 34 of the Freedom of Information and Protection of Privacy Act, for the purposes of determining or verifying eligibility to participate in a program or receive a benefit, product or health service.

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