



PHARMACY COVERAGE GUIDELINES
SECTION: DRUGS

ORIGINAL EFFECTIVE DATE: 9/15/16
LAST REVIEW DATE: 10/19/18
LAST CRITERIA REVISION DATE: 10/19/18
ARCHIVE DATE:

DIRECT ACTING ANTIVIRAL AGENTS FOR HEPATITIS C VIRUS (HCV):

DAKLINZA™ (daclatasvir) oral tablet

EPCLUSA® (velpatasvir, sofosbuvir) oral tablet

HARVONI® (ledipasvir, sofosbuvir) oral tablet

MAVYRET™ (glecaprevir, pibrentasvir) oral tablet

OLYSIO™ (simeprevir sodium) oral capsule

SOVALDI™ (sofosbuvir) oral tablet

TECHNIVIE™ (paritaprevir, ombitasvir, ritonavir) oral tablet

VIEKIRA PAK™ (paritaprevir, ombitasvir, dasabuvir, ritonavir) oral tablet therapy pack

VIEKIRA XR™ (paritaprevir, ombitasvir, dasabuvir, ritonavir) extended release oral tablet

VOSEVI™ (voxilaprevir, velpatasvir, sofosbuvir) oral tablet

ZEPATIER™ (grazoprevir, elbasvir) oral tablet

Coverage for services, procedures, medical devices and drugs are dependent upon benefit eligibility as outlined in the member's specific benefit plan. This Pharmacy Coverage Guideline must be read in its entirety to determine coverage eligibility, if any.

This Pharmacy Coverage Guideline provides information related to coverage determinations only and does not imply that a service or treatment is clinically appropriate or inappropriate. The provider and the member are responsible for all decisions regarding the appropriateness of care. Providers should provide BCBSAZ complete medical rationale when requesting any exceptions to these guidelines.

The section identified as "Description" defines or describes a service, procedure, medical device or drug and is in no way intended as a statement of medical necessity and/or coverage.

The section identified as "Criteria" defines criteria to determine whether a service, procedure, medical device or drug is considered medically necessary or experimental or investigational.

State or federal mandates, e.g., FEP program, may dictate that any drug, device or biological product approved by the U.S. Food and Drug Administration (FDA) may not be considered experimental or investigational and thus the drug, device or biological product may be assessed only on the basis of medical necessity.

Pharmacy Coverage Guidelines are subject to change as new information becomes available.

For purposes of this Pharmacy Coverage Guideline, the terms "experimental" and "investigational" are considered to be interchangeable.

BLUE CROSS®, BLUE SHIELD® and the Cross and Shield Symbols are registered service marks of the Blue Cross and Blue Shield Association, an association of independent Blue Cross and Blue Shield Plans. All other trademarks and service marks contained in this guideline are the property of their respective owners, which are not affiliated with BCBSAZ.

DIRECT ACTING ANTIVIRAL AGENTS FOR HEPATITIS C VIRUS (HCV) (cont.)

This Pharmacy Coverage Guideline does not apply to FEP or other states' Blues Plans.

Information about medications that require precertification is available at www.azblue.com/pharmacy.

Some large (100+) benefit plan groups may customize certain benefits, including adding or deleting precertification requirements.

All applicable benefit plan provisions apply, e.g., waiting periods, limitations, exclusions, waivers and benefit maximums.

Precertification for medication(s) or product(s) indicated in this guideline requires completion of the request form in its entirety with the chart notes as documentation. All requested data must be provided. Once completed the form must be signed by the prescribing provider and faxed back to BCBSAZ Pharmacy Management at (602) 864-3126 or emailed to Pharmacyprecert@azblue.com. **Incomplete forms or forms without the chart notes will be returned.**

Criteria:

- **Criteria for initial therapy:** Daklinza, Epclusa, Harvoni, Mavyret, Olysio, Sovaldi, Technivie, Viekira Pak, Viekira XR, Vosevi, and Zepatier for **treatment Hepatitis C infection with or without ribavirin** is considered **medically necessary** and will be approved when **ALL** of the following criteria are met:
1. Prescriber is a Gastroenterologist, Hepatologist, or Infectious Disease provider
 2. Individual is 18 years of age for Daklinza, Epclusa, Mavyret, Olysio, Technivie, Viekira Pak, Viekira XR, Vosevi, and Zepatier or 12 years of age or older for Harvoni and Sovaldi
 3. A confirmed diagnosis of chronic hepatitis C virus (HCV)
 4. Individual with past hepatitis C treatment has been compliant with previous and/or current drug therapy
 5. There must be no alcohol and/or no substance use in the past 6 months
 6. There must be no contraindications for use of agent(s) requested
 7. There are no significant interacting drugs
 8. For treatment- for HCV for **non-preferred** drugs: Failure, contraindication or intolerance to **at least 2** of the preferred agent by HCV genotype and treatment history
 9. **For HCV treatment requiring concurrent use of ribavirin and the individual is not currently on ribavirin:** Individual has failure, contraindication or intolerance to **generic ribavirin 200mg**
 10. Duration of treatment is consistent with product labeling, current clinical guideline recommendation from AASLD / IDSA for the specific HCV genotype, liver evaluation, individual's treatment status, individual's prior treatment history, and comorbidities

DIRECT ACTING ANTIVIRAL AGENTS FOR HEPATITIS C VIRUS (HCV) (cont.)

- For **Harvoni**: **PREFERRED**
 - Genotype 1: without cirrhosis, HCV RNA level < 6 million IU/mL, HIV-uninfected, non-black: 8 weeks
 - Genotype 1: treatment-naïve, without cirrhosis or compensated cirrhosis, one or more of the following (HCV RNA level \geq 6 million IU/mL, HIV-positive, black): 12 weeks
 - Genotype 1: treatment-experienced, with decompensated cirrhosis (Child-Pugh B & C): 12 weeks with ribavirin
 - Genotype 1: treatment-experienced without cirrhosis: 12 weeks
 - Genotype 1 or 4: Treatment-naïve and treatment-experienced liver transplant recipients without cirrhosis, or with compensated cirrhosis (Child-Pugh class A): 12 weeks with ribavirin
 - Genotype 4,5,6: Treatment-naïve and treatment-experienced without cirrhosis or with compensated cirrhosis (Child-Pugh class A) :12 weeks

- For **Epclusa**: **PREFERRED**
 - Genotypes 1,2,3,4,5,&6: Without cirrhosis or with compensated cirrhosis (Child-Pugh class A):12 weeks
 - Genotypes 1,2,3,4,5,&6: With decompensated cirrhosis (Child-Pugh class B or C): 12 weeks with ribavirin

- For **Mavyret**- **PREFERRED-Without cirrhosis**
 - Genotype 1,2,3,4,5,6: treatment naïve (without cirrhosis): 8 weeks
 - Genotype 1,2,3,4,5,6: treatment naïve (with compensated cirrhosis): 12 weeks
 - Genotype 1 & prior NS3/4A inhibitor but no prior NS5A inhibitor: 12 weeks
 - Genotype 1 & prior NS5A inhibitor but no prior NS3/4A inhibitor: 16 weeks
 - Genotype 3: prior pegylated interferon + ribavirin \pm sofosbuvir (PRS), no prior NS3/4A or NS5A inhibitor: 16 weeks
 - Genotype 1,2,4,5,6: prior PRS, no prior NS3/4A or NS5A inhibitor: 12 weeks
 - Genotype 1,2,4,5,6: no cirrhosis, prior PRS, no prior NS3/4A or NS5A inhibitor: 8 weeks

- For **Vosevi**- **NOT PREFERRED-Without cirrhosis or with compensated cirrhosis (Child-Pugh A)**
 - Genotype 1,2,3,4,5,6: treatment naïve): Prior treatment with NS5A inhibitor:12 weeks
 - Genotype 1a &3: prior treatment with sofosbuvir but without NS5A inhibitor: 12 weeks

- For **Zepatier**-**NOT PREFERRED**
 - Genotype 1a, no NS5A polymorphism, treatment-naïve or prior interferon/ribavirin: 12 weeks
 - Genotype 1a, with NS5A polymorphism, treatment-naïve or prior interferon/ribavirin: 16 weeks
 - Genotype 1b, treatment-naïve or prior interferon/ribavirin: 12 weeks
 - Genotype 1a or b, treatment-naïve or prior interferon/ribavirin/NS3/4A inhibitor: 12 weeks when used with ribavirin
 - Genotype 4, treatment naïve: 12 weeks
 - Genotype 4, prior interferon/ribavirin: 16 weeks

Approval duration: Per HCV genotype and patient specific factors
Prescribing provider must submit viral load after 12 weeks of completion of therapy (SVR12)

DIRECT ACTING ANTIVIRAL AGENTS FOR HEPATITIS C VIRUS (HCV) (cont.)

REQUIRED DOCUMENTATION FOR SUBMISSION OF HCV PRIOR AUTHORIZATION REQUESTS

In order for a prior authorization request for HCV medications to be considered, the following minimum information must be submitted for the member:

1. Evidence of liver fibrosis and presence of cirrhosis plus if it is compensated or decompensated
2. HCV treatment history and responses.
3. Evidence of Hepatitis A & B vaccinations or laboratory evidence of immunity.
4. Current medication list.
5. There must be an abstinence contract signed by the patient and provider
6. An HIV test must be done within 90 days of request for treatment of HCV, if positive, a treatment plan for HCV and HIV therapies must be sent
7. Laboratory results for and drug/alcohol screen completed within the past 90 days.
8. Laboratory results HCV screen, genotype and current baseline viral load within last 90 days
9. Laboratory results for total bilirubin, albumin, INR, CrCl or GFR, LFTs, CBC within past 90 days
10. There must be a negative pregnancy test in a woman of child bearing potential, unless is using adequate contraception

Description:

The presence of hepatitis C (HCV) antibody and HCV RNA are used to support a diagnosis of HCV infection. There are at least six major genotypes and several subtypes of HCV. Baseline viral load by quantitative assay and genotype are necessary to guide therapeutic options.

Hepatitis C infection is a major cause of chronic liver disease and a leading reason for liver transplantation. Sequelae of chronic hepatitis may include liver fibrosis, cirrhosis, liver failure, and hepatocellular carcinoma. Hepatocellular carcinoma rarely progress without underlying fibrosis and cirrhosis.

During acute HCV infection, there is a 20-50% chance of spontaneous resolution of infection. In at least two-thirds of individuals, this will occur within 6 months of the estimated time of infection; only 11% of those who remain viremic at 6 months will spontaneously clear infection at some time later.

Treatment of HCV is rapidly evolving and clinical practice guidelines change as new agents and results of clinical studies become available. Newer agents alone or used in combination with other agents attempt to improve sustained virologic response (SVR) rates, reduce pill-burden, reduce drug-drug interactions, and improve patient tolerance to the medication. The American Association for the Study of Liver Diseases

DIRECT ACTING ANTIVIRAL AGENTS FOR HEPATITIS C VIRUS (HCV) (cont.)

(AASLD) and the Infectious Disease Society of America (IDSA) jointly publish a guideline on the treatment of HCV that can be accessed at <http://www.hcvguidelines.org/full-report-view>. The guideline has recommendations for testing, managing, and treating all HCV genotypes. Treatment options should consider patient-specific factors such as HCV genotype, prior treatment history, presence or absence of compensated or decompensated cirrhosis. The guidance uses evidence-based information.

Ribavirin in combination with an interferon or non-interferon oral anti-hepatitis C antiviral medications is indicated for the treatment of chronic hepatitis C viral (HCV) infection in patients with compensated liver disease. Ribavirin is a synthetic nucleoside analog (purine analog) with antiviral activity. It inhibits replication of RNA and DNA viruses, it inhibits influenza virus RNA polymerase activity and it inhibits the initiation and elongation of RNA fragments resulting in inhibition of viral protein synthesis.

HCV is an RNA virus that utilizes several important enzymes for reproduction. One is a NS3/4A serine protease enzyme that acts to cut large HCV encoded proteins into smaller pieces that are used to build new viruses. It is essential for viral replication. An additional enzyme that is essential for viral replication is NS5B RNA-dependent RNA polymerase that synthesizes the viral genome. The RNA polymerase initiates RNA synthesis by forming a bond between nucleotides that also begins the elongation process of RNA synthesis. A third enzyme, NS5A functions through interaction with other NS viral proteins and other cellular proteins that play a role in mediating host cell function and HCV viral replication, assembly, and egress. Cross-resistance is possible between HCV NS3/4A protease inhibitors and between HCV NS5A inhibitors by class.

About 10-15% of HCV genotype 1 infected patients without prior exposure to NS5A inhibitors will have detectable HCV NS5A resistance associated substitutions (RASs) prior to treatment. The presence of baseline NS5A RASs can cause a large reduction in the activity (> 5 fold) of NS5A inhibitors that potentially adversely impact response to NS5A containing regimens. Given that baseline NS5A RASs are one of the strongest pretreatment predictors of treatment outcome with certain regimens, testing for these RASs prior to deciding on a therapeutic course is now recommended in select situations. Patients with genotype 1a may have higher rates of failure than genotype 1b and RASs testing is recommended for genotype 1a. If the genotype cannot be subtyped recommendations from AASLD is to treat as a genotype 1a infection.

Definitions:

Direct acting antiviral agents for hepatitis C, oral:

NS3/4A serine protease inhibitors:

- Boceprevir – found in Victrelis, no longer available on the market
- Glecaprevir – found in Mavyret
- Grazoprevir – found in Zepatier
- Paritaprevir – found in Viekira Pak, Viekira XR, and Technivie
- Simeprevir – found in Olysio
- Telaprevir – found in Incivek, no longer available on the market
- Voxilaprevir – found in Vosevi

NS5A inhibitors:

- Daclatasvir – found in Daklinza
- Elbasvir – found in Zepatier
- Ledipasvir – found in Harvoni

PHARMACY COVERAGE GUIDELINES
SECTION: DRUGS

ORIGINAL EFFECTIVE DATE: 9/15/16
LAST REVIEW DATE: 10/19/18
LAST CRITERIA REVISION DATE: 10/19/18
ARCHIVE DATE:

DIRECT ACTING ANTIVIRAL AGENTS FOR HEPATITIS C VIRUS (HCV) (cont.)

Ombitasvir – found in Viekira Pak, Viekira XR, and Technivie
Pibrentasvir – found in Mavyret
Velpatasvir – found in Epclusa, Vosevi

NS5B polymerase inhibitors:

Dasabuvir – non-nucleoside inhibitor found in Viekira Pak and Viekira XR
Sofosbuvir – nucleotide inhibitor found in Sovaldi, Harvoni, Epclusa, and Vosevi

CYP3A inhibitors:

Ritonavir – inhibitor of metabolism, found in Viekira Pak, Viekira XR, and Technivie

The Child-Pugh classification system

The Child-Pugh classification is a scoring system used to determine the prognosis of chronic liver disease and cirrhosis. Scoring is based upon several factors: albumin, total bilirubin, prothrombin time or international normalized ratio, and degrees of ascites and encephalopathy

Child-Pugh Classification of severity of liver disease			
Child-Pugh Classification	Points		
A: Well compensated	5-6		
B: Significant functional compromise	7-9		
C: Decompensated	10-15		
Parameter/Factor	1 point each	2 points each	3 points each
Total Bilirubin, mg/dL (or $\mu\text{mol/L}$)	< 2 (or < 34)	2-3 (or 34-50)	>3 or (> 50)
Albumin, g/dL (or g/L)	>3.5 (or > 35)	2.8-3.5 (or 28-35)	< 2.8 (or < 28)
Prothrombin time prolongation			
Seconds over control	1-3	4-6	> 6
INR	< 1.7	1.71-2.3	> 2.3
Ascites	Absent	Slight/Mild	Moderate to severe
Encephalopathy	None	Grade 1-2 (or suppressed with medication)	Grade 3-4 (or refractory)

Hepatitis C Treatment Naive – Preferred Oral Agents

	Genotype 1	Genotype 2	Genotype 3	Genotype 4	Genotype 5	Genotype 6
Epclusa	X	X	X	X	X	X
Harvoni	X			X	X	X
Mavyret†	X	X	X	X	X	X

Harvoni is not FDA approved for genotypes 2,3
† No cirrhosis or compensated cirrhosis (Child-Pugh A)

PHARMACY COVERAGE GUIDELINES
SECTION: DRUGS

ORIGINAL EFFECTIVE DATE: 9/15/16
LAST REVIEW DATE: 10/19/18
LAST CRITERIA REVISION DATE: 10/19/18
ARCHIVE DATE:

DIRECT ACTING ANTIVIRAL AGENTS FOR HEPATITIS C VIRUS (HCV) (cont.)

HCV treatment options by genotype:

FDA-Approved indications						
	Genotype 1	Genotype 2	Genotype 3	Genotype 4	Genotype 5	Genotype 6
Daklinza + Sovaldi ± RBV	X		X			
Epclusa	X	X	X	X	X	X
Harvoni	X			X	X	X
Mavyret†	X	X	X	X	X	X
Olysio* + RBV / Peg-INF	X			X		
Olysio* + Sovaldi	X					
Sovaldi + RBV / Peg-INF	X			X		
Sovaldi + RBV		X	X			
Technivie + RBV				X		
Viekira Pak ± RBV & Viekira XR ± RBV	X					
Vosevi+	X+	X	X+	X	X	X
Zepatier**	X			X		

† It is indicated for treatment-naïve genotype 1-6 patients without cirrhosis or with compensated cirrhosis (C-P Class A) and also indicated for genotype 1 patients without cirrhosis or with compensated cirrhosis (C-P Class A) previously treated for HCV infection with a regimen that contained NS3/4A protease inhibitor or NS5A inhibitor, but not both and for genotypes 1-6 previously treated with pegylated interferon, ribavirin, and/or sofosbuvir, but no prior NS3/4 or NS5A inhibitors

* No Q80K substitution

+ It is indicated for individuals previously treated a HCV regimen: for genotypes 1-6 without cirrhosis or with compensated cirrhosis (C-P Class A) whose previous regimen contained a NS5A inhibitor and for genotype 1a or 3 whose regimen contained sofosbuvir without an NS5A inhibitor

** No baseline NS5A RASs at amino acid positions M28, Q30, L31, and Y93 for genotype 1a

Duration of treatment, depending on regimen, can be as short as weeks 8 weeks but usually ranges 12-24 weeks, and is influenced by genotype, agent(s) selected, prior treatment, presence of compensated or decompensated cirrhosis, and other patient specific factors

RBV = Ribavirin
Peg-INF = pegylated interferon

Ribavirin intolerance or ineligibility – requirements

Platelets < 50,000 cell/mm³

Neutrophils < 750 cell/mm³

Hemoglobin < 10 g/dL

Autoimmune hepatitis or other autoimmune condition known to be exacerbated by ribavirin

**PHARMACY COVERAGE GUIDELINES
SECTION: DRUGS**

**ORIGINAL EFFECTIVE DATE: 9/15/16
LAST REVIEW DATE: 10/19/18
LAST CRITERIA REVISION DATE: 10/19/18
ARCHIVE DATE:**

DIRECT ACTING ANTIVIRAL AGENTS FOR HEPATITIS C VIRUS (HCV) (cont.)

Resources:

Mavyret. Package insert. Revised by manufacturer 8/2017.

Vosevi. Package insert. Revised by manufacturer 7/2017.

Daklinza. Package Insert. Revised by manufacturer 4/2016.

Eplclusa. Package Insert. Revised by manufacturer 6/2016.

Harvoni. Package Insert. Revised by manufacturer 6/2016.

Olysio. Package Insert. Revised by manufacturer 5/2016.

Sovaldi. Package Insert. Revised by manufacturer 8/2015.

Technivie. Package Insert. Revised by manufacturer 6/2016.

Viekira Pak. Package Insert. Revised by manufacturer 6/2016.

Viekira XR. Package Insert. Revised by manufacturer 7/2016.

Rebetol. Package Insert. Revised by manufacturer 1/2016.

Zepatier. Package Insert. Revised by manufacturer 1/2016.

AASLD/IDSA/IAS–USA. Recommendations for Testing, Managing, and Treating Hepatitis C.
<http://www.hcvguidelines.org>. Accessed Oct. 14, 2014

Afdhal N, Reddy KR, Nelson DR et al. Ledipasvir and Sofosbuvir for Previously Treated HCV Genotype 1 Infection. NEJM April 17, 2014 370; 16: 1483-1493

Kowdley KV, Gordon SV, Reddy KR et al. Ledipasvir and Sofosbuvir for 8 or 12 Weeks for Chronic HCV without Cirrhosis NEJM May 15, 2014 370; 19:1879-1888

WHO Guidelines for treatment of hepatitis C <http://www.who.int/mediacentre/factsheets/fs164/en/> Accessed 10/27/14

UpToDate: Overview of the management of chronic hepatitis C virus infection. Current through Aug 2017.
https://www.uptodate-com.mwu.idm.oclc.org/contents/overview-of-the-management-of-chronic-hepatitis-c-virus-infection?source=search_result&search=hepatitis%20c&selectedTitle=1~150

Copegus. Package Insert. Revised by manufacturer 08-2015. Accessed 01-09-2018.

Moderiba. Package Insert. Revised by manufacturer 12-2017. Accessed 01-09-2018.

Rebetol. Package Insert. Revised by manufacturer 01-2016. Accessed 01-09-2018.



PHARMACY COVERAGE GUIDELINES
SECTION: DRUGS

ORIGINAL EFFECTIVE DATE: 9/15/16
LAST REVIEW DATE: 10/19/18
LAST CRITERIA REVISION DATE: 10/19/18
ARCHIVE DATE:

DIRECT ACTING ANTIVIRAL AGENTS FOR HEPATITIS C VIRUS (HCV) (cont.)

Ribasphere. Package Insert. Revised by manufacturer 12-2015. Accessed 07-09-2018.

Ribavirin. Package Insert. Revised by manufacturer 01-2016. Accessed 01-09-2018.

UpToDate: Ribavirin (systemic): Drug information. https://www.uptodate-com.mwu.idm.oclc.org/contents/ribavirin-systemic-drug-information?search=ribavirin&source=search_result&selectedTitle=1~149&usage_type=default&display_rank=1



An Independent Licensee of the Blue Cross and Blue Shield Association

Fax completed prior authorization request form to 602-864-3126 or email to pharmacyprecert@azblue.com. Call 866-325-1794 to check the status of a request. All requested data must be provided. Incomplete forms or forms without the chart notes will be returned. Pharmacy Coverage Guidelines are available at www.azblue.com/pharmacy.

Pharmacy Prior Authorization Request Form

Direct Acting Antiviral Agents for Hepatitis C Virus (HCV)

Do not copy for future use. Forms are updated frequently.

REQUIRED: Office notes, labs, and medical testing relevant to the request that show medical justification are required.

Member Information			
Member Name (first & last):	Date of Birth:	Gender:	BCBSAZ ID#:
Address:	City:	State:	Zip Code:
Prescribing Provider Information			
Provider Name (first & last):	Specialty:	NPI#:	DEA#:
Office Address:	City:	State:	Zip Code:
Office Contact:	Office Phone:	Office Fax:	
Dispensing Pharmacy Information			
Pharmacy Name:	Pharmacy Phone:	Pharmacy Fax:	
Requested Medication Information			
Medication Name:	Strength:	Dosage Form:	
Directions for Use:	Quantity:	Refills:	Duration of Therapy/Use:
<input type="checkbox"/> Check if requesting brand only <input type="checkbox"/> Check if requesting generic			
<input type="checkbox"/> Check if requesting continuation of therapy (prior authorization approved by BCBSAZ expired)			
Preferred agents are Mavyret, Harvoni, and Epclusa			
Turn-Around Time For Review			
<input type="checkbox"/> Standard <input type="checkbox"/> Urgent. Sign here: _____ <input type="checkbox"/> Exigent (requires prescriber to include a written statement)			
Clinical Information			
1. What is the diagnosis? Please specify below. <input type="checkbox"/> Chronic Hepatitis C Virus (HCV) <input type="checkbox"/> Other diagnosis: ICD-10 Code: _____ Diagnosis Description: _____			
2. What is the individual's specific genotype? <input type="checkbox"/> 1 <input type="checkbox"/> 2 <input type="checkbox"/> 3 <input type="checkbox"/> 4 <input type="checkbox"/> 5 <input type="checkbox"/> 6			
3. What is the individual's HCV treatment status? <input type="checkbox"/> Naïve <input type="checkbox"/> Relapse <input type="checkbox"/> Non-responder			
4. What is the prescriber's specialty? <input type="checkbox"/> Gastroenterologist <input type="checkbox"/> Hepatologist <input type="checkbox"/> Infectious Disease <input type="checkbox"/> Other: _____			
5. <input type="checkbox"/> Yes <input type="checkbox"/> No Individual has abstained from alcohol and/or substance use in the past 6 months.			
6. <input type="checkbox"/> Yes <input type="checkbox"/> No <u>For individuals with past HCV treatment:</u> Individual has been compliant with previous and/or current drug therapy.			
7. <input type="checkbox"/> Yes <input type="checkbox"/> No <u>For HCV treatment requiring concurrent use of ribavirin:</u> Individual has failure, contraindication or intolerance to generic ribavirin 200mg.			
8. <input type="checkbox"/> Yes <input type="checkbox"/> No There is absence of all contraindications for use of agent(s) requested.			
9. <input type="checkbox"/> Yes <input type="checkbox"/> No There is absence of all significant interacting drugs.			
10. <input type="checkbox"/> Yes <input type="checkbox"/> No <u>Required documentation:</u> Evidence of liver fibrosis and presence of cirrhosis; plus, if it is compensated or decompensated.			
11. <input type="checkbox"/> Yes <input type="checkbox"/> No <u>Required documentation:</u> HCV treatment history and responses.			
12. <input type="checkbox"/> Yes <input type="checkbox"/> No <u>Required documentation:</u> Child-Pugh score - total bilirubin, albumin, INR within past 90 days plus status of ascites and encephalopathy.			
13. <input type="checkbox"/> Yes <input type="checkbox"/> No <u>Required documentation:</u> Evidence of Hepatitis A & B vaccinations or laboratory evidence of immunity.			

Pharmacy Prior Authorization Request Form

Direct Acting Antiviral Agents for Hepatitis C Virus (HCV)

14. <input type="checkbox"/> Yes <input type="checkbox"/> No	Required documentation: <u>Current medication list.</u>
15. <input type="checkbox"/> Yes <input type="checkbox"/> No	Required documentation: <u>Abstinence contract signed by the patient and provider.</u>
16. <input type="checkbox"/> Yes <input type="checkbox"/> No	Required documentation: <u>An HIV test must be done within 90 days of request for treatment of HCV; if positive, a treatment plan for HCV and HIV therapies must be sent.</u>
17. <input type="checkbox"/> Yes <input type="checkbox"/> No	Required documentation: <u>Laboratory results for and drug/alcohol screen completed within the past 90 days.</u>
18. <input type="checkbox"/> Yes <input type="checkbox"/> No	Required documentation: <u>Laboratory results HCV screen, genotype and current baseline viral load within last 90 days.</u>
19. <input type="checkbox"/> Yes <input type="checkbox"/> No	Required documentation: <u>Laboratory results for CrCl or GFR, LFTs, CBC within past 90 days.</u>
20. <input type="checkbox"/> Yes <input type="checkbox"/> No	Required documentation (if applicable): <u>A negative pregnancy test in a woman of child bearing potential, unless is using adequate contraception.</u>

21. What medication(s) has the individual tried and failed for this diagnosis? Please specify below.

Important note: Samples provided by the provider are not accepted as continuation of therapy or as an adequate trial and failure.

Medication Name, Strength, Frequency	Dates started and stopped or Approximate Duration	Describe response, reason for failure, or allergy

22. Are there any supporting labs or test results? Please specify below.

Date	Test	Value

23. Is there any additional information the prescribing provider feels is important to this review? Please specify below.

For example, explain the negative impact on medical condition, safety issue, reason formulary agent is not suitable to a specific medical condition, expected adverse clinical outcome from use of formulary agent, or reason different dosage form or dose is needed.

Signature affirms that information given on this form is true and accurate and reflects office notes	
Prescribing Provider's Signature:	Date:

Please note: Some medications may require completion of a drug-specific request form.

Incomplete forms or forms without the chart notes will be returned.

Office notes, labs, and medical testing relevant to the request that show medical justification are required.