



Original Effective Date: 01/01/2022  
Current Effective Date: 01/01/2026  
Last P&T Approval/Version: 10/29/2025  
Next Review Due By: 10/2026  
Policy Number: C21903-A

## Hepatitis C Antiviral Therapy

### PRODUCTS AFFECTED

Epclusa (sofosbuvir/velpatasvir), Harvoni (ledipasvir/sofosbuvir), ledipasvir-sofosbuvir, Mavyret (glecaprevir and pibrentasvir), ribavirin, sofosbuvir-velpatasvir, Sovaldi (sofosbuvir), Viekira Pak (paritaprevir/ritonavir/ombitasvir and dasabuvir), Vosevi (sofosbuvir, velpatasvir, voxilaprevir), Zepatier (elbasvir and grazoprevir)

### COVERAGE POLICY

*Coverage for services, procedures, medical devices, and drugs are dependent upon benefit eligibility as outlined in the member's specific benefit plan. This Coverage Guideline must be read in its entirety to determine coverage eligibility, if any. This 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 Molina Healthcare complete medical rationale when requesting any exceptions to these guidelines.*

#### **Documentation Requirements:**

*Molina Healthcare reserves the right to require that additional documentation be made available as part of its coverage determination; quality improvement; and fraud; waste and abuse prevention processes. Documentation required may include, but is not limited to, patient records, test results and credentials of the provider ordering or performing a drug or service. Molina Healthcare may deny reimbursement or take additional appropriate action if the documentation provided does not support the initial determination that the drugs or services were medically necessary, not investigational, or experimental, and otherwise within the scope of benefits afforded to the member, and/or the documentation demonstrates a pattern of billing or other practice that is inappropriate or excessive.*

#### **DIAGNOSIS:**

Chronic Hepatitis C Infection

#### **REQUIRED MEDICAL INFORMATION:**

This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved. If a drug within this policy receives an updated FDA label within the last 180 days, medical necessity for the member will be reviewed using the updated FDA label information along with state and federal requirements, benefit being administered and formulary preferencing. Coverage will be determined on a case-by-case basis until the criteria can be updated through Molina Healthcare, Inc. clinical governance. Additional information may be required on a case-by-case basis to allow for adequate review. When the requested drug product for coverage is dosed by weight, body surface area or other member specific measurement, this data element is required as part of the medical necessity review. The Pharmacy and Therapeutics Committee has determined that the drug benefit shall be a mandatory generic and that generic drugs will be dispensed whenever available.

#### **A. FOR ALL INDICATIONS:**

##### **1. Documentation of diagnosis of Hepatitis C virus (HCV) Infection**

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AND

2. Documentation requested medication is prescribed in accordance with the current FDA approved labeling and current AASLD guideline recommendation at the dose and duration appropriate for the member

AND

3. For chronic infection: Documentation of HCV RNA (HCV viral load)

OR

For acute infection (Mavyret only): Documentation of HCV RNA (HCV viral load) and negative HCV antibody within the past 6 weeks

AND

4. Documentation of HIV status (if positive see HIV CO- INFECTION requirements), and cirrhosis status (e.g., Child Pugh Class, FIB-4 score, FibroScan, FibroSure, clinical evidence of cirrhosis, prior liver biopsy, etc.) [DOCUMENTATION REQUIRED]

AND

5. FOR REGIMENS THAT INCLUDE RIBAVIRIN OR AASLD REGIMENS THAN RECOMMEND RIBAVIRIN:

(a) Documentation member or member's partner is not pregnant, two reliable forms of contraception will be used during therapy, and pregnancy tests will be performed as needed throughout treatment AND Documentation of CBC within last 6 months AND Prescriber attests member has been counseled on preventing pregnancy for up to 6 months after stopping regimen AND

(b) For exception to FDA approved regimens that contain ribavirin for Ribavirin-ineligible members, documentation of FDA labeled contraindication to Ribavirin is required: pregnancy, CrCl <30 ml/minute, Hepatic decompensation (Child-Pugh class B and C), Hemoglobin <8.5 g/dL, WBC <1,000 mm<sup>3</sup>, neutrophils <500 mm<sup>3</sup> or Platelets <50 x 10<sup>9</sup>/L

AND

6. Documentation of prior HCV treatment history and if treatment-experienced, include date(s) of all HCV therapy (start and completion) and the resulting outcome or discontinuation status and reason for discontinuation.

AND

7. Documentation of the following Resistant Associated Substitutions (RASs) testing: REQUIRED ONLY FOR REQUESTS WITH THE APPLICABLE DRUG REGIMEN AND MEMBER/TREATMENT CHARACTERISTICS DETAILED BELOW:

- a. Zepatier (elbasvir/grazoprevir): NS5A RAS testing is recommended for genotype 1a-infected, treatment-naïve or -experienced members being considered for elbasvir/grazoprevir
- b. Harvoni (sofosbuvir/ledipasvir): NS5A RAS testing can be considered for genotype 1a-infected, treatment-experienced patients with and without cirrhosis being considered for ledipasvir/sofosbuvir. If >100-fold resistance is present, a different recommended therapy should be used
- c. Epclusa (sofosbuvir/velpatasvir): NS5A RAS testing is recommended for genotype 3-infected, treatment-experienced patients with or without cirrhosis and treatment-naïve patients with cirrhosis being considered for 12 weeks of sofosbuvir/velpatasvir. If Y93H is present, weight-based ribavirin should be added or another recommended regimen should be used. (AASLD I, A)
- d. Daklinza (daclatasvir) plus Sovaldi (sofosbuvir): NS5A RAS testing is recommended for genotype 3-infected, treatment-experienced patients without cirrhosis being considered for 12 weeks of daclatasvir plus sofosbuvir. Indicate if baseline NS5A Y93H polymorphism is present. If Y93H is present, weight-based ribavirin should be added. (AASLD I, B) NS5A RAS testing is recommended for genotype 3-infected, treatment-naïve patients with compensated cirrhosis being considered for 24 weeks of daclatasvir plus sofosbuvir. Indicate if baseline NS5A Y93H polymorphism is present. If Y93H is present, treatment should include weight-based ribavirin, or a different therapy should be used. (AASLD I, B)

**MOLINA REVIEWER NOTE:** If genotype testing is not captured in another requirement, genotype

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testing is required for these regimen requests when the above member/treatment characteristics are met to review RAS testing per AASLD guidelines.

AND

8. FOR NON-FORMULARY/NON-PREFERRED AGENTS: Documented rationale of clinical medical necessity for the inability to utilize the FORMULARY/PREFERRED AGENTS
- B. CHRONIC HEPATITIS C INFECTION TREATMENT, RE-TREATMENT, RE-INFECTION:
  1. Documentation of HCV genotype and subtype (obtained within the last 3 years and for current infection if re-infection) with a confirmed genotype of 1a, 1b, 2, 3, 4, 5, or 6 [DOCUMENTATION REQUIRED]  
AND
  2. (a) Documentation of reason(s) for previous failure of DAA therapy (i.e., tolerability, efficacy, etc.) and if the stated reason(s) for the previous treatment failure have been addressed (e.g., re-education and understanding/agreement of treatment plan)  
OR  
(b) Documentation (post therapy sustained virologic response [SVR] lab) member was successfully treated with DAA therapy and has subsequently been re-infected.  
NOTE: Inadequate compliance to regimen and treatment plan (including office visits, scheduled clinical tests, etc.) or non-adherence to initial/previous HCV regimen as evidenced by medical records and/or pharmacy claims should be noted and addressed  
AND
  3. Member adherent to previous HCV therapy as evidenced by pharmacy claims [MOLINA REVIEWER VERIFY]  
AND
  4. Documentation of ONE of the following:
    - (a) Documentation of failure to achieve a sustained virologic response (SVR) or lack of efficacy during treatment as evidenced by detectable serum HCV RNA by quantitative assay at 12 or more weeks after completing treatment; or a 10-fold increase of viral load at week 6 of treatment.  
[DOCUMENTATION REQUIRED: Laboratory documentation of quantitative viral load required.]  
OR  
(b) Documentation of adverse event that required therapy discontinuation as evidenced by laboratory results (e.g., CBC, LFTs, etc.) and/or clinical presentation, AND No improvement of adverse effect after proper clinical management  
OR  
(c) Documentation of re-infection (i.e., report of high-risk behavior since previous treatment)  
AND
    5. Requested regimen is the highest-rated regimen per AASLD Guidelines for member's condition [by viral subtype, previous therapy, presence or absence of cirrhosis, and presence or absence of resistance-associated variants (RAVs)]  
AND
    6. There is evidence that such re-treatment will improve member outcomes according to AASLD guidelines OR at least a Class IIa rating (weight of evidence and/or opinion is in favor of usefulness and efficacy) or higher per AASLD Guidelines

## C. CHRONIC HEPATITIS C INFECTION, PEDIATRICS

1. (a) Documentation of HCV genotype and subtype (obtained within the last 3 years) with a confirmed genotype of 1a, 1b, 2, 3, 4, 5, or 6 [DOCUMENTATION REQUIRED]  
OR  
(b) Request is for an AASLD regimen for children or adolescents recommended with any genotype  
AND
2. Requested regimen must meet the American Association for the Study of Liver Diseases and the Infectious Diseases Society of America Present HCV Guidance: Recommendations for Testing, Managing, and Treating Hepatitis: HCV in Children AND the most recent FDA labeled indications

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**D. CHRONIC HEPATITIS C INFECTION, HIV COINFECTION:**

1. Documentation of diagnosis of HIV-1  
AND
2. Documentation of HCV genotype and subtype (obtained within the last 3 years) with a confirmed genotype of 1a, 1b, 2, 3, 4, 5, or 6 [DOCUMENTATION REQUIRED]

**E. DECOMPENSATED CIRRHOSIS (CHILD PUGH B OR C):**

1. Documentation of Decompensated liver disease  
AND
2. Documentation of HCV genotype and subtype (obtained within the last 3 years) with a confirmed genotype of 1a, 1b, 2, 3, 4, 5, or 6 [DOCUMENTATION REQUIRED]

**F. HEPATOCELLULAR CARCINOMA AWAITING LIVER TRANSPLANT:**

1. Documentation of diagnosis of Stage I-III hepatocellular carcinoma (HCC) confirmed by image testing (ultrasound, tomography, MRI, laparoscopy), or biopsy  
NOTE: It is reasonable to treat HCV in a member with HCC or a history of HCC after the HCC has been treated successfully, with follow-up imaging demonstrating locoregional control. Patients with HCC should be assessed for DAA therapy on a case-by-case basis and, ideally, managed with input from a Tumor Board or specialty care. Patients with extensive or progressive HCC (e.g., vascular invasion or metastatic disease) are less likely to benefit from DAA therapy  
AND
2. Member meets all criteria for authorization of a liver transplant as indicated in Molina Healthcare MCP- 114: Liver Transplantation Adult & Pediatric  
AND
3. Documentation of HCV genotype and subtype (obtained within the last 3 years) with a confirmed genotype of 1a, 1b, 2, 3, 4, 5, or 6 [DOCUMENTATION REQUIRED]

### **CONTINUATION OF THERAPY:**

For new to Molina Members, authorization should be entered to allow completion of regimen up to appropriate AASLD duration - 8,12,16, or 24 weeks.

For re-treatment - Review as new authorization

Acute treatment – NA

### **DURATION OF APPROVAL:**

Per appropriate AASLD regimen 8, 12, 16, or 24 weeks

### **PRESCRIBER REQUIREMENTS:**

**ACUTE OR CHRONIC HEPATITIS C INFECTION, TREATMENT NAÏVE, WITHOUT CIRRHOSIS:** No prescriber

Requirement

**HEPATITIS C INFECTION, HIV COINFECTION:** Prescribed by or in consultation with an infectious disease specialist

**ALL OTHER INDICATIONS:** Prescribed by or in consultation with a gastroenterologist, hepatologist, infectious disease, or transplant medicine specialist

[If prescribed in consultation, consultation notes must be submitted with the initial request.]

### **AGE RESTRICTIONS:**

Acute infection Mavyret only: 3 years of age and older

18 years of age and older with the exceptions found under CHRONIC HEPATITIS C INFECTION, PEDIATRICS section

### **QUANTITY:**

Max quantity is a 28 day supply per dispense. The number of dispenses are allowed up to the approved

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duration.

### PLACE OF ADMINISTRATION:

The recommendation is that oral medications in this policy will be for pharmacy benefit coverage and patient self-administered.

## DRUG INFORMATION

### ROUTE OF ADMINISTRATION:

Oral

### DRUG CLASS:

Hepatitis C Agents

### FDA-APPROVED USES:

*EPCLUSA (sofosbuvir/velpatasvir)* is indicated for the treatment of adult patients and pediatric patients 3 years of age and older with chronic HCV genotype 1, 2, 3, 4, 5, or 6 infection: without cirrhosis or with compensated cirrhosis OR with decompensated cirrhosis for use in combination with ribavirin.

*HARVONI (ledipasvir/sofosbuvir)* is indicated for the treatment of chronic hepatitis C virus (HCV) in adults and pediatric patients 3 years of age and older with genotype 1, 4, 5, or 6 infection without cirrhosis or with compensated cirrhosis, genotype 1 infection with decompensated cirrhosis, in combination with ribavirin, genotype 1 or 4 infection who are liver transplant recipients without cirrhosis or with compensated cirrhosis, in combination with ribavirin.

*MAVYRET (glecaprevir and pibrentasvir)* is indicated for the treatment of adult and pediatric patients 3 years and older with acute or chronic HCV genotype (GT) 1, 2, 3, 4, 5 or 6 infection without cirrhosis or with compensated cirrhosis (Child-Pugh A) AND indicated for the treatment of adult and pediatric patients 3 years and older with HCV genotype 1 infection, who previously have been treated with a regimen containing an HCV NS5A inhibitor or an NS3/4A protease inhibitor, but not both.

*Ribavirin*: indicated in combination with interferon alfa-2b (pegylated and nonpegylated) for the treatment of Chronic Hepatitis C (CHC) in patients 3 years of age or older with compensated liver disease. Patients with the following characteristics are less likely to benefit from re-treatment after failing a course of therapy: previous nonresponse, previous pegylated interferon treatment, significant bridging fibrosis or cirrhosis, and genotype 1 infection.

*SOVALDI (sofosbuvir)* is indicated for the treatment of adult patients with genotype 1, 2, 3 or 4 chronic hepatitis C virus (HCV) infection without cirrhosis or with compensated cirrhosis as a component of a combination antiviral treatment regimen AND pediatric patients 3 years of age and older with genotype 2 or 3 chronic HCV infection without cirrhosis or with compensated cirrhosis in combination with ribavirin.

*VIEKIRA PAK (paritaprevir/ritonavir/ombitasvir and dasabuvir)* is indicated for the treatment of adult patients with chronic hepatitis C virus (HCV) with genotype 1b without cirrhosis or with compensated cirrhosis AND genotype 1a without cirrhosis or with compensated cirrhosis for use in combination with ribavirin.

*VOSEVI (sofosbuvir, velpatasvir, voxilaprevir)* is indicated for the treatment of adult patients with chronic HCV infection without cirrhosis or with compensated cirrhosis (Child-Pugh A) who have genotype 1, 2, 3, 4, 5, or 6 infection and have previously been treated with an HCV regimen containing an NS5A inhibitor AND genotype 1a or 3 infection and have previously been treated with an HCV regimen containing sofosbuvir without an NS5A inhibitor. Additional benefit of Vosevi over sofosbuvir/velpatasvir was not shown in adults with genotype 1b, 2, 4, 5, or 6 infection previously.

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treated with sofosbuvir without an NS5A inhibitor.

*ZEPATIER (elbasvir and grazoprevir tablet)* is indicated for treatment of chronic HCV genotype 1 or 4 infection in adult and pediatric patients 12 years of age and older or weighing at least 30 kg. Zepatier is indicated for use with ribavirin in certain patient populations.

## COMPENDIAL APPROVED OFF-LABELED USES:

None

## APPENDIX

### APPENDIX:

#### **AASLD recommendations for patients co-infected with HIV and HCV**

HIV/HCV-coinfected patients should be treated and retreated the same as patients without HIV infection, after recognizing and managing interactions with antiretroviral medications (AASLD Class I, Level B). Antiretroviral drug switches, when needed, should be done in collaboration with HIV practitioner; for HIV antiretroviral and HCV direct-acting antiviral combinations not addressed within the guideline, expert consultation is recommended (AASLD Class I, Level A)

## BACKGROUND AND OTHER CONSIDERATIONS

### BACKGROUND:

Risk of Hepatitis B infection reactivation with HCV Direct Acting Antivirals (DAA) In October of 2016, the FDA issued a safety alert concerning risk of reactivation of hepatitis B viral (HBV) infection in patients treated with HCV direct acting antivirals (DAA). At the time of the alert, the FDA had identified 24 cases of HBV infection reactivation in patients who had been treated with a HCV DAA. In a few of these cases, the HBV reactivation resulted in serious liver problems or death. As a result, the FDA has required labeling for all HCV DAAAs to include a boxed warning for HBV infection reactivation. In addition, the FDA has recommended that all patients be screened for evidence of current or prior HBV infection before starting treatment with HCV DAAAs and be monitored for HBV reactivation during and after treatment with a HCV DAA.

### CONTRAINDICATIONS/EXCLUSIONS/DISCONTINUATION:

All other uses of direct acting antivirals are considered experimental/investigational and therefore, will follow Molina's Off- Label policy.

Contraindications to Epclusa (sofosbuvir/velpatasvir) include: Epclusa and ribavirin combination regimen is contraindicated in patients for whom ribavirin is contraindicated.

Contraindications to Harvoni (ledipasvir/sofosbuvir) include: If used in combination with ribavirin, all contraindications to ribavirin also apply to Harvoni combination therapy.

Contraindications to Mavyret (glecaprevir and pibrentasvir) include: Patients with moderate or severe hepatic impairment (Child-Pugh B or C) or those with any history of prior hepatic decompensation, coadministration with atazanavir or rifampin.

Contraindications to Ribavirin include: pregnancy and men whose female partners are pregnant, known hypersensitivity reactions such as Stevens-Johnson syndrome, toxic, epidermal necrolysis, and erythema multiforme to ribavirin or any component of the product, autoimmune hepatitis, hemoglobinopathies, creatinine clearance less than 50 mL/min, coadministration with didanosine.

Contraindications to Sovaldi (sofosbuvir) include: when used in combination with peginterferon alfa/ribavirin or ribavirin alone, all contraindications to peginterferon alfa and/or ribavirin also apply to Sovaldi

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combination therapy.

Contraindications to Viekira Pak (paritaprevir/ritonavir/ombitasvir and dasabuvir) include: patients with moderate to severe hepatic impairment, If VIEKIRA PAK is administered with ribavirin, the contraindications to ribavirin also apply to this combination regimen, co-administration with drugs that are: highly dependent on CYP3A for clearance; moderate or strong inducers of CYP3A and strong inducers of CYP2C8; and strong inhibitors of CYP2C8, known hypersensitivity to ritonavir (e.g. toxic epidermal necrolysis, Stevens-Johnson syndrome).

Contraindications to Vosevi (sofosbuvir, velpatasvir, voxilaprevir) include: coadministration with rifampin.

Contraindications to Zepatier (elbasvir and grazoprevir) include: Patients with moderate or severe hepatic impairment (Child-Pugh B or C), OATP1B1/3 inhibitors that are known or expected to significantly increase grazoprevir plasma concentrations, strong CYP3A inducers, and efavirenz, If Zepatier is administered with ribavirin, the contraindications to ribavirin also apply.

**Treatment is recommended for all patients with acute or chronic HCV infection, except those with a short life expectancy that cannot be remediated by HCV therapy, liver transplantation, or another directed therapy (AASLD, May 2023). Patients with a short life expectancy owing to liver disease should be managed in consultation with an expert**

Members identified having any barriers to treatment mentioned in RMI are not appropriate candidates for therapy until issues have been resolved, or acknowledgement of actions taken by prescriber or another provider involved in the member's care to address those barriers

Pregnancy: For women of reproductive age with known HCV infection, antiviral therapy is recommended before considering pregnancy, whenever practical and feasible, to reduce risk of HCV transmission to future offspring. (AASLD, October 2022).

The safety and efficacy of DAA therapy in pregnant or lactating women have not been established for any of the currently FDA-approved agents. During pregnancy, these drugs should be used only if the benefits outweigh the risks to the fetus.

Severe end organ disease and is not eligible for solid organ transplant. Clinically significant illness or any other major medical disorder that may interfere with a patient's ability to complete a course of treatment. Individuals who in the professional judgment of the primary treating clinician would not achieve a long-term clinical benefit from HCV treatment, with conditions such as those: Multisystem organ failure, Receiving palliative care or are enrolled in hospice, Presence of significant pulmonary or cardiac disease, Malignancy outside of the liver not meeting oncologic criteria for cure, Decompensated liver disease with CTP score > 12 or MELD > 20, OR Model For End-Stage Liver Disease (MELD) ≤ 20 and ONE (1) of the following: [ONE] Cardiopulmonary disease that cannot be correct and is a prohibitive risk for surgery, Malignancy outside of the liver not meeting oncologic criteria for cure, Hepatocellular carcinoma with metastatic spread or not listed for liver transplant, Intrahepatic cholangiocarcinoma, Hemangiosarcoma Decompensated liver disease with CTP score > 12 or MELD > 20, OR Model For End-Stage Liver Disease (MELD) ≤ 20 and ONE (1) of the following: [ONE] Cardiopulmonary disease that cannot be correct and is a prohibitive risk for surgery, Malignancy outside of the liver not meeting oncologic criteria for cure, Hepatocellular carcinoma with metastatic spread or not listed for liver transplant, Intrahepatic cholangiocarcinoma , Hemangiosarcoma

## OTHER SPECIAL CONSIDERATIONS:

The AASLD/IDSA 2023 updated guidance for testing, managing, and treating Hepatitis C includes a new recommendation that addresses the management of incomplete treatment adherence. The algorithm for the

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management of incomplete adherence as part of DAA treatment monitoring is applicable only to DAA treatment-naïve persons and, generally, the same patient populations who are eligible for the simplified treatment algorithms. Excluded persons with incomplete adherence should be managed in consultation with a specialist in HCV management.

Recommended management of DAA treatment interruptions for treatment-naïve patients without cirrhosis or with compensated cirrhosis receiving glecaprevir/pibrentasvir or sofosbuvir/velpatasvir (Figure 2, Bhattacharya et al, 2023)

### Interruptions BEFORE Receiving 28 Days of DAA Therapy:

- Missed ≤7 days
  - Restart DAA therapy immediately. Complete therapy for originally planned duration (8 or 12 weeks).
- Missed ≥8 days
  - Restart DAA therapy immediately. Restarting DAA takes precedence over obtaining HCV RNA level.
  - Obtain HCV RNA test as soon as possible, preferably the same day as restarting the DAA therapy.
    - See guideline for specific recommendations based on HCV RNA outcome.

### Interruptions AFTER Receiving ≥28 Days of DAA Therapy

- Missed ≤7 days
  - Restart DAA therapy immediately. Complete therapy for originally planned duration (8 or 12 weeks).
- Missed 8-20 Consecutive Days
  - Restart DAA therapy immediately. Restarting DAA takes precedence over obtaining HCV RNA level.
  - Obtain HCV RNA test as soon as possible, preferably the same day as restarting the DAA therapy.
    - See guideline for specific recommendations based on HCV RNA outcome.
- Missed ≥21 Consecutive Days
  - Stop DAA treatment and assess for sVR12. If SVR12 not achieved, retreat according to recommendations in the Retreatment Section.

Epclusa, Mavyret, Zepatier, Harvoni, Sovaldi, Vosevi, Viekira Pak have a black box warning for risk of hepatitis B virus reactivation in patients coinfected with HCV and HBV. Hepatitis B virus (HBV) reactivation has been reported, in some cases resulting in fulminant hepatitis, hepatic failure, and death.

Ribavirin has a black box warning for embryo-fetal toxicity, hemolytic anemia, and monotherapy not recommended.

## CODING/BILLING INFORMATION

**CODING DISCLAIMER.** Codes listed in this policy are for reference purposes only and may not be all-inclusive or applicable for every state or line of business. Deleted codes and codes which are not effective at the time the service is rendered may not be eligible for reimbursement. Listing of a service or device code in this policy does not guarantee coverage. Coverage is determined by the benefit document. Molina adheres to Current Procedural Terminology (CPT®), a registered trademark of the American Medical Association (AMA). All CPT codes and descriptions are copyrighted by the AMA; this information is included for informational purposes only. Providers and facilities are expected to utilize industry-standard coding practices for all submissions. Molina has the right to reject/deny the claim and recover claim payment(s) if it is determined it is not billed appropriately or not a covered benefit. Molina reserves the right to revise this policy as needed.

HCPCS CODE	DESCRIPTION
NA	

## AVAILABLE DOSAGE FORMS:

Epclusa PACK 150-37.5MG

Epclusa PACK 200-50MG

Epclusa TABS 200-50MG

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Epclusa TABS 400-100MG  
Harvoni PACK 33.75-150MG  
Harvoni PACK 45-200MG  
Harvoni TABS 45-200MG  
Harvoni TABS 90-400MG  
Ledipasvir-Sofosbuvir TABS 90-400MG  
Mavyret PACK 50-20MG  
Mavyret TABS 100-40MG  
Ribavirin CAPS 200MG  
Ribavirin TABS 200MG  
Sofosbuvir-Velpatasvir TABS 400-100MG  
Sovaldi PACK 150MG  
Sovaldi PACK 200MG Sovaldi TABS 200MG  
Sovaldi TABS 400MG  
Viekira Pak TBPK 12.5-75-50 &250MG  
Vosevi TABS 400-100-100MG  
Zepatier TABS 50-100MG

## REFERENCES

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2. Harvoni (ledipasvir/sofosbuvir) [prescribing information]. Foster City, CA: Gilead Sciences Inc; March 2020.
3. Mavyret (glecaprevir/pibrentasvir) [prescribing information]. North Chicago, IL: AbbVie Inc; June 2025.
4. Ribavirin [prescribing information]. East Windsor, NJ: Aurobindo Pharma USA, Inc.; July 2023.
5. Sovaldi (sofosbuvir) [prescribing information]. Foster City, CA: Gilead Sciences; March 2020.
6. Viekira Pak (ombitasvir/paritaprevir/ritonavir/dasabuvir) [prescribing information]. North Chicago, IL: AbbVie Inc; December 2019.
7. Vosevi (sofosbuvir, velpatasvir, voxilaprevir) [prescribing information]. Foster City, CA: Gilead Sciences, Inc; November 2019.
8. Zepatier (elbasvir and grazoprevir) tablets [prescribing information]. Rahway, NJ: Merch Sharp & Dohme LLC; May 2022.
9. AASLD-IDSA. Recommendations for testing, managing, and treating hepatitis C. <http://www.hcvguidelines.org>. [31 October 2022].
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12. American Association for the Study of Liver Diseases AASLD and Infectious Diseases Society of America IDSA. (2022, October 24). Management of Acute HCV Infection | HCV Guidance. Retrieved from www.hcvguidelines.org website: <https://www.hcvguidelines.org/unique-populations/acute-infection>

## Drug and Biologic Coverage Criteria

SUMMARY OF REVIEW/REVISIONS	DATE
REVISION- Notable revisions: Required Medical Information Prescriber Requirements Age Restrictions FDA-Approved Uses References	Q4 2025
REVISION- Notable revisions: Required Medical Information Prescriber Requirements Quantity Contraindications/Exclusions/Discontinuation	Q1 2025
REVISION- Notable revisions: Products Affected Required Medical Information FDA-Approved Uses Contraindications/Exclusions/Discontinuation Other Special Considerations Available Dosage Forms References	Q1 2024
REVISION- Notable revisions: Products Affected Required Medical Information Prescriber Requirements FDA- Approved Uses Appendix Contraindications/Exclusions/Discontinuation References	Q1 2023
Q2 2022 Established tracking in new format	Historical changes on file