

PHARMACY POLICY STATEMENT

HAP CareSource™ Marketplace

| DRUG NAME | Sofosbuvir/Velpatasvir (Epclusa) |
|--------------|----------------------------------|
| BENEFIT TYPE | Pharmacy |
| STATUS | Prior Authorization Required |

Sofosbuvir/Velpatasvir (Epclusa) is indicated for the treatment of adults 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. It is also indicated for the treatment of adults and pediatric patients 3 years of age and older with chronic HCV genotype 1, 2, 3, 4, 5, or 6 infection with decompensated cirrhosis for use in combination with ribavirin.

Sofosbuvir is a HCV nucleotide analog NS5B polymerase inhibitor that prevents hepatitis C viral replication through RNA chain termination. Velpatasvir prevents viral replication through inhibition of NS5A protein.

Sofosbuvir/Velpatasvir (Epclusa) will be considered for coverage when the following criteria are met:

Hepatitis C (without cirrhosis or with compensated cirrhosis (Child-Turcotte-Pugh Class A))



For **initial** authorization:

- 1. Member must be 3 years of age or older; AND
- 2. Member has a diagnosis of hepatitis C without cirrhosis or with compensated cirrhosis (Child-Turcotte-Pugh Class A); AND
- 3. Member has genotype 1, 2, 3, 4, 5 or 6 (laboratory documentation required); AND
- 4. Member's documented viral load taken within 6 months of beginning therapy and submitted with chart notes.
- 5. Dosage allowed/Quantity limit:
 - a) Adult patients: Take one tablet (400 mg of sofosbuvir and 100 mg of velpatasvir) once daily for 12 weeks.
 - b) Pediatric patients 3 years and older:

| Body weight (kg) | Epclusa Daily Dose | Dosing of Epclusa Oral Pellets | Dosing of Epclusa Tablet |
|-----------------------|----------------------|---|-----------------------------------|
| Less than 17 kg | 150mg/37.5mg per day | One 150mg/37.5mg packet of pellets once daily | N/A |
| 17 to less than 30 kg | 200mg/50mg per day | One 200mg/50mg packet of pellets once daily | One 200mg/50mg tablet once daily |
| At least 30 kg | 400mg/100mg per day | Two 200mg/50mg packets of pellets once daily | One 400mg/100mg tablet once daily |

If all the above requirements are met, the medication will be approved for 12 weeks.

For reauthorization:

1. Medication will not be reauthorized.

Hepatitis C with Decompensated Cirrhosis (Child-Turcotte-Pugh Class B or C)

For **initial** authorization:

- 1. Member must be 3 years of age or older; AND
- 2. Member has a diagnosis of hepatitis C with decompensated cirrhosis (Child-Turcotte-Pugh Class B or C); AND
- 3. Member has genotype 1, 2, 3, 4, 5 or 6 (laboratory documentation required); AND
- 4. Provider attests member will be prescribed sofosbuvir/velpatasvir (generic for Epclusa) in combination with ribavirin when applicable; AND
- 5. Member's documented viral load taken within 6 months of beginning therapy and submitted with chart notes.
- 6. Dosage allowed/Quantity limit:

<u>Adult patients</u>: Take one tablet (400 mg of sofosbuvir and 100 mg of velpatasvir) once daily for 12 weeks. If member is ribavirin ineligible, sofosbuvir/velpatasvir (generic for Epclusa) may be approved for a total of 24 weeks.

Pediatric patients 3 years of age or older:

| Body weight (kg) | Epclusa Daily Dose | Dosing of Epclusa Oral | Dosing of Epclusa |
|------------------|---------------------------|------------------------|-------------------|
| | | Pellets | Tablet |



| Less than 17 kg | 150mg/37.5mg per day | One 150mg/37.5mg packet | N/A |
|--------------------|----------------------|-------------------------|-------------------|
| - | | of pellets once daily | |
| 17 to less than 30 | 200mg/50mg per day | One 200mg/50mg packet | One 200mg/50mg |
| kg | | of pellets once daily | tablet once daily |
| At least 30 kg | 400mg/100mg per day | Two 200mg/50mg packets | One 400mg/100mg |
| | | of pellets once daily | tablet once daily |

If all the above requirements are met, the medication will be approved for 12 weeks for ribavirin eligible members. If the request is for a ribavirin ineligible member, the medication will be approved for 24 weeks.

For reauthorization:

1. Medication will not be reauthorized.

HAP CareSource considers Sofosbuvir/Velpatasvir (Epclusa) not medically necessary for the treatment of conditions that are not listed in this document. For any other indication, please refer to the Off-Label policy.

| DATE | ACTION/DESCRIPTION | |
|------------|--|--|
| 05/09/2017 | New policy for Epclusa created | |
| 06/08/2017 | Fibrosis stage 2 and above covered. | |
| 11/22/2017 | Medication status changed to non-preferred. Substance abuse program information is no longer required. Trial of preferred agent is required for members without cirrhosis or with compensated cirrhosis only | |
| 12/07/2017 | Criterion of "life expectancy not less than one year due to non-liver related comorbidities" removed from criteria and added in a form of the note. Hepatitis B testing is no longer required. | |
| 12/21/2017 | Fibrosis score requirement was removed. | |
| 05/01/2019 | Policy modified to Sofosbuvir/velpatasvir (generic for Epclusa); status changed to preferred product. Trial of Mavyret removed. | |
| 04/26/2020 | Age requirement criterion changed from 18 years old to 6 years old or weighing 17 kg (37 lbs) for both diagnoses. | |
| 11/18/2021 | Updated age requirement to 3 years and older; Updated reference section; Transferred to new policy template | |
| 02/24/2023 | Removed drug screen requirement. Updated pediatric dosing information. | |
| 04/12/2023 | Removed prescriber specialty requirement. | |
| 11/14/2023 | Updated/added/removed references. Changed genotype requirement from 1,2,3,4,6 to 1,2,3,4,5,6 for patients who have decompensated cirrhosis and are ribavirin ineligible; Decreased initial approval duration from 12 months to up to 24 weeks for decompensated cirrhosis and 12 weeks for no cirrhosis/compensated cirrhosis; Removed specific documentation proving member is ribavirin ineligible (hemoglobin, neutrophils, platelets etc). | |
| 03/27/2024 | Added pediatric dosing to Hepatitis C (without cirrhosis or with compensated cirrhosis) section; specified adult dosing. | |



04/10/2025

Changed title from Epclusa to Sofosbuvir/Velpatasvir (Epclusa)

04/22/2025

Updated references; removed note "Member's life expectancy must be no less than one year due to non-liver related comorbidities" to align with other hepatitis C policies; added provider attestation to ribavirin combination use and removed note about submitting documentation of clinical reason ribavirin cannot be used; added confirmation of hepatitis C diagnosis

References:

- 1. Epclusa [package insert]. Foster City, CA: Gilead Sciences Inc.; 2022.
- 2. Bhattacharya D, Aronsohn A, Price J, Lo Re V; AASLD-IDSA HCV Guidance Panel . Hepatitis C Guidance 2023 Update: AASLD-IDSA Recommendations for Testing, Managing, and Treating Hepatitis C Virus Infection [published online ahead of print, 2023 May 25]. *Clin Infect Dis*. 2023;ciad319. doi:10.1093/cid/ciad319
- 3. Clinical Care of Hepatitis C. Centers for Disease Control and Prevention. Updated January 31, 2025. Accessed April 22, 2025. https://www.cdc.gov/hepatitis-c/hcp/clinical-care/index.html

Effective date: 10/01/2025 Revised date: 04/22/2025