

Hepatitis C: Sovaldi

Member Information

1. Last Name: _____ 2. First Name: _____
3. Trillium ID #: _____ 4. Date of Birth: _____ 5. Gender: _____

Prescriber Information

1. Prescriber Name: _____ 2. NPI #: _____
3. Requestor Name (Nurse/Office Staff): _____
4. Mailing Address: _____ City: _____ State: _____ Zip: _____
5. Phone #: _____ Ext. _____ Fax #: _____

Drug Information

1. Drug Name: Sovaldi 2. Strength: _____ 3. Quantity per 30 Days: 28
4. Length of Therapy (in days): 365 days

Clinical Information

Total Length of Therapy (Check ONE):

- 12 weeks = Genotype 1, 2, or 4 for treatment-naïve and treatment-experienced adult beneficiaries without cirrhosis or with compensated cirrhosis (child-pugh A); or genotype 2 for treatment-naïve and treatment-experienced pediatric patients, 3 years of age or older, without cirrhosis or with compensated cirrhosis (child-pugh A). Genotype 1 and previously treated with a regimen containing an NS3/4A PI2 without prior treatment with an NS5A inhibitor.
 - 24 weeks = Genotype 1 adult beneficiaries that are PEG-interferon ineligible; genotype 3 for treatment-naïve and treatment-experienced adults without cirrhosis or with compensated cirrhosis (child-pugh A); or genotype 3 for treatment-naïve and treatment-experienced pediatric patients, 3 years of age or older, without cirrhosis or with compensated cirrhosis (child-pugh A)
 - 48 weeks = Genotype 1,2,3, or 4 for adult beneficiaries with a diagnosis of Hepatocellular Carcinoma awaiting liver transplantation (up to 48 weeks or until liver transplantation, whichever comes first)
1. What is the member's Genotype? _____
 2. Is the member 18 years of age or older with a diagnosis of Chronic Hepatitis C infection with confirmed genotype 1 or 4 without cirrhosis or with compensated cirrhosis? Yes No
 3. Is the member 3 years of age or older with a diagnosis of Chronic Hepatitis C infection with confirmed genotype 2 or 3 without cirrhosis or with compensated cirrhosis? Yes No
 4. Does the member have a CHC infection with hepatocellular carcinoma awaiting a liver transplant?
 Yes No
 5. As the provider, are you reasonably certain that treatment will improve the member's overall health status? Yes No
 6. Is Sovaldi being prescribed in combination with: Ribavirin and pegylated Interferon alfa for Genotype 1 or 4 Ribavirin for beneficiaries with genotype 1 who are peginterferon-ineligible (medical record documentation of previous peginterferon therapy or reason for ineligibility must be submitted for review) Ribavirin for Genotypes 2 or 3 and/or in CHC patients with hepatocellular carcinoma awaiting liver transplant
 7. Is Sovaldi being used as monotherapy? Yes No
 8. Is Sovaldi being used with any other sofosbuvir containing regimen? Yes No
 9. Does the member have FDA labeled contraindication to sofosbuvir (Sovaldi)? Yes No
 10. Is the member pregnant? Yes No
 11. Does the member have severe renal impairment (CrCl less than 30mL/min), end stage renal disease, or requires dialysis (AASLD/IDSA 2014)? Yes No
 12. Is the member a non-responder to sofosbuvir? Yes No
 13. Has the member previously failed therapy with a treatment regimen that included sofosbuvir? Yes No
 14. Does the member have hepatocellular carcinoma and is not awaiting liver transplant? Yes No

Fax this form to PerformRx at (833) 726-7628 or call Pharmacy PA Call Center: (855) 662-0277

Signature of Prescriber: _____ Date: _____

(Prescriber Signature Mandatory)

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.