

# Sovaldi™ Initiation Prior Authorization Form

Member Name: \_\_\_\_\_ Date of Birth: \_\_\_\_\_ Member ID#: \_\_\_\_\_  
 Pharmacy NPI: \_\_\_\_\_ Pharmacy Phone: \_\_\_\_\_ Pharmacy Fax: \_\_\_\_\_  
 Pharmacy Name: \_\_\_\_\_ Pharmacist Name: \_\_\_\_\_  
 Prescriber NPI: \_\_\_\_\_ Prescriber Name: \_\_\_\_\_ Specialty: \_\_\_\_\_  
 Prescriber Phone: \_\_\_\_\_ Prescriber Fax: \_\_\_\_\_ Drug Name: \_\_\_\_\_  
 NDC: \_\_\_\_\_ Start Date: \_\_\_\_\_

## Clinical Information

- Diagnosis: \_\_\_\_\_ HCV Genotype (including subtype): \_\_\_\_\_
- METAVIR Fibrosis Stage: \_\_\_\_\_ Date Determined: \_\_\_\_\_
- Does member have decompensated hepatic disease (CTP class B or C)? Yes \_\_\_ No \_\_\_
- Has the member been evaluated by a gastroenterologist, infectious disease specialist, or a transplant specialist within the past 3 months? Yes \_\_\_ No \_\_\_
- If yes, please include name of specialist recommending hepatitis C treatment: \_\_\_\_\_
- Please indicate regimen below (if member is IFN ineligible please specify reasoning):
  - Sovaldi™ 400mg daily w/ weight-based RBV plus weekly PEG/IFN x 84 days (12 weeks)
  - Sovaldi™ 400mg daily w/ weight-based RBV x 84 days (12 weeks)
  - Sovaldi™ 400mg daily w/ weight-based RBV x 168 days (24 weeks)
  - Other: \_\_\_\_\_ \*\*
  - Sovaldi™ 400mg daily w/ weight-based RBV x 336 days (48 weeks)\*\*  
 \*\*Approvals for 48 weeks will only be granted for HCV infected members with hepatocellular carcinoma meeting the MI-LAN criteria. Does member have tumor ≤5cm in diameter with single hepatocellular carcinomas and not more than 3 tumor nodules, each ≤3cm in diameter and no extrahepatic manifestations of the cancer or evidence of vascular invasion of tumor? Yes \_\_\_ No \_\_\_
- Has the member signed the intent to treat contract\*\*? Yes \_\_\_ No \_\_\_  
 \*\*Required for processing of prior authorization request
- Has the member had illicit IV drug use or alcohol abuse in the last 6 months? Yes \_\_\_ No \_\_\_
- Has the member initiated immunization with the hepatitis A and B vaccines? Yes \_\_\_ No \_\_\_
- For women of childbearing potential (and male patients with female partners of childbearing potential):
  - Patient is not pregnant (or a male with a pregnant female partner) and not planning to become pregnant during treatment or within 6 months of completing treatment
  - Agreement that partners will use two forms of effective non-hormonal contraception during treatment and for at least 6 months after completing treatment  
Please list non-hormonal birth control options discussed with member \_\_\_\_\_
  - Verification that monthly pregnancy tests will be performed throughout treatment
- Is the member taking any of the following medications: amiodarone, rifampin, rifabutin, rifapentine, carbamazepine, eslicarbazepine, phenytoin, oxcarbazepine, tipranavir/ritonavir, didanosine or St. John's wort? Yes \_\_\_ No \_\_\_
- Have all other clinically significant issues been addressed prior to starting therapy? Yes \_\_\_ No \_\_\_
- I recommend this patient be followed by an OHCA Care Management Nurse.

**Members must be adherent for continued approval. Treatment gaps of therapy longer than 3 days will result in denial of payment for subsequent requests for continued therapy. Refills must be prior authorized.**

Prescriber Signature: \_\_\_\_\_ Date: \_\_\_\_\_  
 Has the member been counseled on appropriate use of Sovaldi™ therapy? Yes \_\_\_ No \_\_\_

Pharmacist Signature: \_\_\_\_\_ Date: \_\_\_\_\_  
 Please do not send in chart notes. Specific information/documentation will be requested if necessary.

PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

University of Oklahoma College of Pharmacy  
Pharmacy Management Consultants  
Product Based Prior Authorization Unit

Fax: 1-800-224-4014  
Phone: 1-800-522-0114 Option 4

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