

**Harvoni® (Ledipasvir/Sofosbuvir) 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: \_\_\_\_\_ Start Date: \_\_\_\_\_  
 Drug Name: \_\_\_\_\_ NDC: \_\_\_\_\_ Member's Weight (kg): \_\_\_\_\_ Date Taken: \_\_\_\_\_

**Clinical Information**

1. HCV Genotype (including subtype): \_\_\_\_\_ Date Determined: \_\_\_\_\_
2. METAVIR Equivalent Fibrosis Stage: \_\_\_\_\_ Testing Type: \_\_\_\_\_  
Date Fibrosis Stage Determined: \_\_\_\_\_
3. Pre-treatment viral load in the last 12 months (must be within last 3 months if requesting 8-week regimen):  
Pre-treatment viral load: \_\_\_\_\_ Date Taken: \_\_\_\_\_  
For METAVIR score of <F1, 2nd test must confirm chronic HCV diagnosis at least 6 months after 1st test.  
Prior pre-treatment viral load or antibody test: \_\_\_\_\_ Date Taken: \_\_\_\_\_
4. Does member have decompensated hepatic disease (CTP class B or C)? Yes \_\_\_\_\_ No \_\_\_\_\_
5. Is the member currently on hospice or does the member have a limited life expectancy (less than 12 months) that cannot be remediated by treating HCV? Yes \_\_\_\_\_ No \_\_\_\_\_
6. Has the member been evaluated by a gastroenterologist, infectious disease specialist, or a transplant specialist within the past 3 months? Yes \_\_\_\_\_ No \_\_\_\_\_
7. If yes, please include name of specialist recommending hepatitis C treatment: \_\_\_\_\_
8. Has the member been previously treated for hepatitis C? Yes \_\_\_\_\_ No \_\_\_\_\_
9. If yes, please indicate previous treatment regimen and reason for failure (relapser, null-responder, partial responder): \_\_\_\_\_
10. Please indicate requested drug strength **and** regimen below:
 

<input type="checkbox"/> Harvoni® 90mg/400mg	<input type="checkbox"/> once daily x 56 days (8 weeks)
<input type="checkbox"/> Harvoni® 45mg/200mg	<input type="checkbox"/> once daily x 84 days (12 weeks)
<input type="checkbox"/> Harvoni® 33.75mg/150mg	<input type="checkbox"/> once daily with weight-based ribavirin x 84 days (12 weeks)
<input type="checkbox"/> Other: _____	
11. For members 6 years of age or older requesting the oral pellet formulation, please provide a patient-specific, clinically significant reason why the tablet is not appropriate: \_\_\_\_\_
12. Has the member signed the intent to treat contract\*\*? Yes \_\_\_\_\_ No \_\_\_\_\_ *\*\*Required for processing of request \*\**
13. Has the member been counseled on the harms of illicit IV drug use and alcohol use and agreed to not use illicit IV drugs or alcohol while on or after they finish hepatitis C treatment? Yes \_\_\_\_\_ No \_\_\_\_\_
14. Has the member initiated immunization with the hepatitis A and B vaccines? Yes \_\_\_\_\_ No \_\_\_\_\_
15. For women of childbearing potential (and male patients with female partners of childbearing potential):
 

<input type="checkbox"/> Patient is not pregnant (or a male with a pregnant female partner) and not planning to become pregnant during treatment
<input type="checkbox"/> Agreement that partners will use 2 forms of effective non-hormonal contraception during treatment (and for 6 months after therapy completion for those on ribavirin). Please list non-hormonal birth control options discussed with member _____
16. Is the member taking any of the following medications: amiodarone, rifampin, rifabutin, rifapentine, carbamazepine, eslicarbazepine, phenytoin, phenobarbital, oxcarbazepine, tipranavir/ritonavir, simeprevir, rosuvastatin, St. John's wort, or elvitegravir/cobicistat/emtricitabine in combination with tenofovir disoproxil fumarate?  
Yes \_\_\_\_\_ No \_\_\_\_\_
17. Have all other clinically significant issues been addressed prior to starting therapy? Yes \_\_\_\_\_ No \_\_\_\_\_

**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 Harvoni® therapy? Yes \_\_\_\_\_ No \_\_\_\_\_

**Pharmacist Signature:** \_\_\_\_\_ **Date:** \_\_\_\_\_

*Please do not send in chart notes. Specific information/documentation will be requested if necessary. Failure to complete this form in full will result in processing delays. By signature, the prescriber or pharmacist confirms the above information is accurate.*

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Fax completed prior authorization request form to 888-601-8461 or submit Electronic Prior Authorization through CoverMyMeds® or SureScripts. All requested data must be provided. Incomplete forms or forms without the chart notes will be returned. Pharmacy Coverage Guidelines are available at [AetnaBetterHealth.com/Oklahoma](http://AetnaBetterHealth.com/Oklahoma).