

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 if applicable): _____ Date Determined: _____
2. METAVIR Equivalent Fibrosis Stage: _____ Testing Type: _____
Date Fibrosis Stage Determined: _____
3. Pre-treatment viral load in the last 12 months: _____ 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 regimen below:
 - sofosbuvir/velpatasvir 400mg/100mg daily x 84 days (12 weeks)
 - sofosbuvir/velpatasvir 400mg/100mg daily with weight-based ribavirin x 84 days (12 weeks)
 - sofosbuvir/velpatasvir 200mg/50mg daily x 84 days (12 weeks)
 - sofosbuvir/velpatasvir 200mg/50mg daily with weight-based ribavirin x 84 days (12 weeks)
 - Other: _____
11. Has the member signed the intent to treat contract**? Yes ___ No ___ ***Required for processing of request*
12. 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 ___
13. Has the member initiated immunization with the hepatitis A and B vaccines? Yes ___ No ___
14. 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
 - 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 _____
15. Is the member taking any of the following medications: H2-receptor antagonists at doses greater than 40mg famotidine equivalent, amiodarone, omeprazole or other proton pump inhibitors, topotecan, rifampin, rifabutin, rifapentine, carbamazepine, eslicarbazepine, phenytoin, phenobarbital, oxcarbazepine, efavirenz, tenofovir disoproxil fumarate, tipranavir/ritonavir, St. John's wort, or rosuvastatin doses exceeding 10mg? Yes ___ No ___
16. If member is using antacids have they agreed to separate antacid and sofosbuvir/velpatasvir administration by 4 hours? Yes ___ No ___ NA ___
17. Have all other clinically significant issues been addressed prior to starting therapy? Yes ___ No ___

This patient is in need of additional support. 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 sofosbuvir/velpatasvir therapy? Yes ___ No ___

Pharmacist Signature: _____ **Date:** _____

Please do not send in chart notes. Failure to complete this form in full will result in processing delays. By signature, the prescriber or pharmacist confirms the above information is accurate.

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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