

Viekira Pak™ (Ombitasvir/Paritaprevir/Ritonavir/Dasabuvir) 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

1. Diagnosis: _____ HCV Genotype (including subtype): _____
 2. METAVIR Fibrosis Stage: _____ Date Determined: _____
 3. Pre-Treatment Viral Load (HCV RNA): _____ Date Determined: _____
 4. Does member have decompensated hepatic disease (CTP class B or C)? Yes ___ No ___
 5. Has the member been evaluated by a gastroenterologist, infectious disease specialist, or a transplant Specialist for hepatitis C therapy within the past 3 months? Yes ___ No ___
 6. If yes, please include name of specialist recommending hepatitis C treatment: _____
 7. Has the member been previously treated for hepatitis C? Yes ___ No ___
 8. If yes, please indicate previous treatment regimen and reason for failure: _____
9. Please indicate requested regimen below:
- Genotype 1a, without cirrhosis: Viekira Pak™ with weight-based ribavirin x 84 days (12 weeks)
 - Genotype 1a with cirrhosis: Viekira Pak™ with weight-based ribavirin x 168 days (24 weeks)
 - Genotype 1b without cirrhosis: Viekira Pak™ x 84 days (12 weeks)
 - Genotype 1b with cirrhosis: Viekira Pak™ with weight-based ribavirin X 84 days (12 weeks)
 - Other: _____ **
- **Please supply reference citation to support requested therapy.*
10. Has the member signed the intent to treat contract**? Yes ___ No ___
***Required for processing of prior authorization request*
 11. Has the member had illicit IV drug use or alcohol abuse in the last 6 months? Yes ___ No ___
 12. Has the member initiated immunization with the hepatitis A and B vaccines? Yes ___ No ___
 13. 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 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
 14. Is the member taking any of the following medications: alfuzosin, carbamazepine, phenytoin, phenobarbital, gemfibrozil, rifampin, ergotamine, dihydroergotamine, ergonovine, methylergonovine, ethinyl estradiol, St. John's wort, lovastatin, simvastatin, pimozone, efavirenz, sildenafil, triazolam, midazolam? Yes ___ No ___
 15. Have all other clinically significant issues been addressed prior to starting therapy? Yes ___ No ___
 16. Will the member's ALT levels be monitored during the first four weeks of starting treatment and as clinically indicated thereafter? 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 Viekira Pak™ 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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