

## Hepatitis C Prior Authorization Request Form (Page 1 of 2)

|  |   |   | DATED FREQUENTLY AN  | ND MAY BE B    | BARCODED        |                |
|--|---|---|--|----------------|-----------------|----------------|
| Memb   | er Information  | (required)  | Provide  | r Inforn       | nation (r       | equired)       |
| Member Name:   |   |   | Provider Name:   |                |                 |                |
| Insurance ID#:   |   |   | NPI#: Specialty:   |                |                 |                |
| Date of Birth:   |   |   | Office Phone:  |                |                 |                |
| Street Address:  |   |   | Office Fax:  |                |                 |                |
| City:  | State:  | Zip:  | Office Street Address:                                       |                |                 |                |
| Phone:   |   |   | City:  | State:         |                 | Zip:           |
|  | Ν   | ledication Info   | mation (required)  |                |                 |                |
| Medication Name:   |   |   | Strength: Dosage Form:                                       |                |                 |                |
| Check if requesting brand  |   |   | Directions for Use:  |                |                 |                |
| Check if request is  | for <b>continuation of the</b>                            | rapy  |  |                |                 |                |
|  |   |   | otion  |                |                 |                |
| Select the diagnosis   | below:  | Clinical Inform   | ation (required)   |                |                 |                |
| Hepatitis C virus in   |   |   |  |                |                 |                |
| <ul> <li>Other diagnosis:</li> </ul>                                 |   |   | ICD-10 Code(s):  |                |                 |                |
| Clinical information:  | ·····   |   |  |                |                 |                |
|  | s genotype:   |   |  |                |                 |                |
|  | s weight:   |   |  |                |                 |                |
| -  |   | oviders may be asked to                                 | provide documentation  | ):             |                 |                |
|  | ve cirrhosis? D Yes                                       |   | F  | ,-             |                 |                |
|  |   | isease (Child-Pugh A)?                                  |  |                |                 |                |
|  |   | r disease (Child-Pugh B                                 | or C)? 🛛 Yes 🖾 No  |                |                 |                |
| •  | nt naïve? 🛛 Yes 🖾 No                                      |   |  |                |                 |                |
| Select one of the follow   | •   |   |  |                |                 |                |
|  | rin intolerant/ineligible                                 |   |  |                |                 | <b>C U U</b>   |
|  |   | in, patient has a negative                              | e pregnancy test within t                                    | 30 days prioi  | r to initiation | of therapy and |
|  | onthly pregnancy test d                                   | uring treatment   |  |                |                 |                |
| Patient is not pr  |   | n aliaibla  |  |                |                 |                |
| -  | ribed ribavirin or ribaviri                               | -   |  |                |                 |                |
|  | -   | svir, also answer the fo                                | -  |                |                 |                |
|  |   | n sofosbuvir or NS5A-ba                                 |  |                |                 |                |
|  |   | ucers (e.g., rifampin, St.                              | John's wort)?  | NO             |                 |                |
|  | nticancers (e.g., topotec                                 | ,   |  |                |                 |                |
| Is the patient taking m oxcarbazepine)?                              |   | nducers (e.g., rifampin, s                              | St. John's wort, carbama                                     | azepine, phe   | enytoin, pher   | iobarbital,    |
|  |   | elpatasvir) in combinatior<br>Zepatier (elbasvir/grazop |  | ct acting anti | viral agent [   | e.g., Sovaldi  |
| For Harvoni or gener   | ric ledipasvir/sofosbuv                                   | vir, also answer the foll                               | owing:   |                |                 |                |
|  | taking any of the followi                                 |   |  |                |                 |                |
|  |   | xcarbazepine, phenobar                                  | bital, phenytoin)  |                |                 |                |
|  | ²-gp) inducers (e.g., rifa<br>s (e.g., tipranavir/ritonav |   |  |                |                 |                |
| <ul> <li>Hiv antiretroviration</li> <li>Tenofovir-contair</li> </ul> |   | /11 /   |  |                |                 |                |
| Anticaners (e.g.,  |   |   |  |                |                 |                |
| Is the patient receiving   |   |   | ith another HCV direct acting antiviral agent [e.g., Sovaldi |                |                 |                |
|  | . /3  |   |  |                |                 |                |

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## Hepatitis C Prior Authorization Request Form (Page 2 of 2) DO NOT COPY FOR FUTURE USE. FORMS ARE UPDATED FREQUENTLY AND MAY BE BARCODED

| For Mavyret (glecaprevir-pibrentasvir), a<br>Select if the patient has been previously tre             | ated with a regimen containing the following (select all that applies):  |  |  |  |  |
|--|--|--|--|--|--|
| □ An HCV NS5A inhibitor  |  |  |  |  |  |
| An NS3/4A protease inhibitor (PI)  |  |  |  |  |  |
| Interferon (including pegylated formula  | ations), ribavirin, and/or Sovaldi (sofosbuvir)  |  |  |  |  |
| Is the patient receiving Mayvret in combina<br>(ledipasvir/sofosbuvir), Zepatier (elbasvir/gi          | tion with another HCV direct acting antiviral agent [e.g., Sovaldi (sofosbuvir), Harvoni razoprevir)]? <b>D Yes D No</b> |  |  |  |  |
| For Sovaldi (sofosbuvir), also answer th   | e following:   |  |  |  |  |
| Select if the patient will use Sovaldi in coml<br>Pegylated interferon and ribavirin<br>Ribavirin      | bination with the following:   |  |  |  |  |
| Does the patient have severe renal impairn   | nent (eGFR < mL/min/1.73 m²)? □ Yes □ No   |  |  |  |  |
| Does the patient have end-stage renal dise   | ase? 🛛 Yes 🖾 No  |  |  |  |  |
| Does the patient have hepatocellular carcinoma that meets criteria for liver transplant? <b>Yes No</b> |  |  |  |  |  |
| For Vosevi (sofosbuvir-velpatasvir-voxil   | aprevir), also answer the following:   |  |  |  |  |
| Has the patient been previously treated with   | h a regimen containing an NS5A inhibitor? 🛛 Yes 🗳 No   |  |  |  |  |
| Has the patient been previously treated with   | h a regimen containing Sovaldi (sofosbuvir) without an NS5A inhibitor? 🛛 Yes 🗳 No  |  |  |  |  |
| Is the patient receiving Vosevi in combination (ledipasvir/sofosbuvir), Zepatier (elbasvir/generation) | on with another HCV direct acting antiviral agent [e.g., Sovaldi (sofosbuvir), Harvoni<br>razoprevir)]? 🛛 Yes 🛛 No       |  |  |  |  |
| For Zepatier (elbasvir-grazoprevir), also  | answer the following:  |  |  |  |  |
| Has the patient been tested for the presence   | ce of NS5A resistance-associated polymorphisms?  Yes  No   |  |  |  |  |
| Does the patient have moderate to severe   | hepatic impairment? 🛛 Yes 🗳 No   |  |  |  |  |
| Is the patient receiving Zepatier in combina (ledipasvir/sofosbuvir)]? <b>☐ Yes ☐ No</b>               | tion with another HCV direct acting antiviral agent [e.g., Sovaldi (sofosbuvir), Harvoni                                 |  |  |  |  |

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note:

This request may be denied unless all required information is received. For urgent or expedited requests please call 1-855-401-4262. This form may be used for non-urgent requests and faxed to 1-844-403-1029.