



## PRIOR AUTHORIZATION REQUEST: Hepatitis C Treatment

*Please fax form to Amida Care: 1-646-786-0997*

### MEMBER INFORMATION

Name:	Medicaid ID #:
Phone #:	Address:

### PRESCRIBER INFORMATION

Name:	NPI:
Office Phone #:	Office Fax #:
Address:	
Contact Person:	

**Please Choose One or More Prescriber Specialty:**

Prescriber Specialty:  Hepatologist  Gastroenterologist  Transplant  Infectious Disease  Other \_\_\_\_\_

### FORMULARY PREFERRED OPTIONS

Select	Indication	Presence of Cirrhosis	Medication Requested	Duration
<input type="checkbox"/>	Genotype 1, 2, 3, 4, 5, or 6: <b>Treatment Naïve</b>	Without cirrhosis	glecaprevir/pibrentasvir (Mavyret®)	8 weeks
<input type="checkbox"/>	Genotype 1, 2, 3, 4, 5, or 6: <b>Treatment Naïve</b>	Without cirrhosis or compensated cirrhosis	velpatasvir/sofosbuvir (ag Eplclusa*)	12 weeks
<input type="checkbox"/>	Genotype 1, 2, 3, 4, 5, or 6: <b>Treatment Naïve</b>	With decompensated cirrhosis	velpatasvir/sofosbuvir (ag Eplclusa*) + Ribavirin	12 weeks
<input type="checkbox"/>	Genotype 1, 2, 4, 5, or 6: <b>Treatment experienced with IFN + RBV +/- sofosbuvir</b>	Without cirrhosis	glecaprevir/pibrentasvir (Mavyret®)	8 weeks
<input type="checkbox"/>	Genotype 1, 2, 3, 4, 5, or 6: <b>Treatment experienced with IFN + RBV +/- sofosbuvir</b>	Without cirrhosis	velpatasvir/sofosbuvir (ag Eplclusa*)	12 weeks
<input type="checkbox"/>	Genotype 1, 2, 4, 5, or 6: <b>Treatment experienced with IFN + RBV +/- sofosbuvir</b>	With decompensated cirrhosis	velpatasvir/sofosbuvir (ag Eplclusa*)	12 weeks
<input type="checkbox"/>	Genotype 3: <b>Treatment experienced with IFN + RBV +/- sofosbuvir</b>	With decompensated cirrhosis	velpatasvir/sofosbuvir (ag Eplclusa*) + Ribavirin	12 weeks
<input type="checkbox"/>	Genotype 1, 2, 3, 4, 5, or 6: <b>Treatment experienced with IFN + RBV +/- sofosbuvir</b>	With decompensated cirrhosis	velpatasvir/sofosbuvir (ag Eplclusa*) + Ribavirin	12 weeks
<input type="checkbox"/>	Genotype 1: <b>Treatment experienced with NS3/4A PI without prior NS5A inhibitor</b>	Without cirrhosis	velpatasvir/sofosbuvir (ag Eplclusa*)	12 weeks
<input type="checkbox"/>	Genotype 1: <b>Treatment experienced with NS3/4A PI without prior NS5A inhibitor</b>	Without cirrhosis or compensated cirrhosis	velpatasvir/sofosbuvir (ag Eplclusa*) + Ribavirin	12 weeks
<input type="checkbox"/>	Genotype 1: <b>Treatment experienced with NS5A inhibitor without prior NS3/4A PI</b>	Without cirrhosis or with compensated cirrhosis	glecaprevir/pibrentasvir (Mavyret®)	16 weeks

*\*The authorized generic form of Eplclusa (velpatasvir/sofosbuvir) is the preferred formulary agent*

**Clinical Rationale required for non-preferred (non-formulary) drug with documentation of contraindication to Eplclusa or Mavyret:**

**Treatment Readiness:** (documentation required)

- Patient demonstration of readiness, willingness, and ability to adhere to the regimen

**Education Readiness:** (documentation required)

- Patient understands reinfection of Hepatitis C is still possible after being cured of Hepatitis C
- Patient understands not to engage in risky and unhealthy behaviors which would lead to reinfection.

**\*\*Amida Care resources are available to support member adherence and lifestyle modification. Please check below to request any additional type of support or services for member:**

- Additional support needed for member by Amida Care (Please specify type of support or education needed):

\_\_\_\_\_

\_\_\_\_\_

**MEDICAL DIAGNOSIS AND CLINICAL CRITERIA**

**Please provide labs/documentation required for verification of questions**

1. Hep C Genotype: \_\_\_\_\_
2. Most recent Baseline HCV RNA Viral titer/Viral Load (within 3 months): \_\_\_\_\_
3. What is the patient's prior treatment status?
- Treatment Naïve     Prior Relapse     Prior Partial Responder     Null Responder     Reinfection

PRIOR HEP C TREATMENT	DURATION/YEAR	OUTCOME OF TREATMENT
	/	
	/	
	/	

**4. Please provide documentation of most recent lab values for CD4 and HIV viral load. (Check appropriate box.)**

- Patient on ARVs and recent viral load under 200 copies/mL with CD4 count above 200
- Patient electing not to be on ARVs and CD4 above 500
- Neither (**Please provide explanation and documentation**)

**5. Please provide documentation of Hepatitis B virus (HBV) status:**

- HBV negative     Concurrent HBV is being treated     Neither (**Please provide explanation and documentation**)

**6. Has the patient had a liver enzyme level test and either a liver biopsy, Fibrosan or Fibrosure diagnostic test?**

*Please provide documentation demonstrating liver enzyme levels and stage of liver fibrosis or cirrhosis.*

- YES     NO    **If cirrhosis is present please check:**     Child Pugh     Child Pugh B     Child Pugh C

**7. Does the prescriber agree to submit HCV-RNA levels at 4 weeks, end of treatment and 3 months post treatment (12 week SVR)?**

- YES     NO    *If NO, please explain:* \_\_\_\_\_

**Please call 646-757-7615 or email [dgreenidge@amidacareny.org](mailto:dgreenidge@amidacareny.org) M-F, 9:00-4:30 PM with questions or additional info. You may also provide us with your contact information and the best time to reach you in the space at the top of this document.**

\_\_\_\_\_  
Prescriber or Authorized Signature

\_\_\_\_\_  
Date