

OPTIMA HEALTH PLAN

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

Directions: The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; **fax to 1-800-750-9692**. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. Incomplete form will delay authorization process.

Hepatitis-C Antiviral Drugs (Non-Preferred) (FAMIS Only)

DRUG REQUESTED – check the applicable box below for Hepatitis-C therapy:			
<input type="checkbox"/> Daklinza™	<input type="checkbox"/> Epclusa®	<input type="checkbox"/> Harvoni®	<input type="checkbox"/> Olysio™
<input type="checkbox"/> Sovaldi®	<input type="checkbox"/> Technivie™	<input type="checkbox"/> Viekira Pak™	<input type="checkbox"/> Viekira XR™
<input type="checkbox"/> Vosevi®	<input type="checkbox"/> Zepatier®	<input type="checkbox"/> peginterferon alfa	<input type="checkbox"/> ribavirin
<input type="checkbox"/> sofosbuvir (400mg)/ velpatasvir (100mg)	<input type="checkbox"/> ledipasvir (90mg)/ sofosbuvir (400mg)		

DRUG INFORMATION: Complete information below or authorization will be delayed.

Drug Name: _____

Drug Form/Strength: _____ **Quantity per Day:** _____

Dosing Schedule: _____ **Length of Therapy:** _____

Diagnosis: _____ **ICD Code, if applicable:** _____

- Optima Health coverage criteria for the new direct-acting agents are based on careful consideration of the evidence-based guidance of professional specialty societies, published guidelines, and physician subject matter experts specialists.
- **ONE TIME APPROVAL FOR ANY and ALL DIRECT-ACTING ANTIVIRAL (DAA) PER LIFETIME (EXCEPTION)**

CLINICAL CRITERIA: Check **ALL** boxes below to qualify. **ALL** pertinent chart notes and lab values **MUST** be included in this request or authorization process will be delayed. Check applicable box(es) below for **PREVIOUS** Hepatitis-C treatments/therapy:

<input type="checkbox"/> Daklinza™	<input type="checkbox"/> Epclusa®	<input type="checkbox"/> Harvoni®	<input type="checkbox"/> Incivek®
<input type="checkbox"/> Olysio™	<input type="checkbox"/> Sovaldi®	<input type="checkbox"/> Technivie™	<input type="checkbox"/> Victrelis®
<input type="checkbox"/> Viekira Pak™	<input type="checkbox"/> Viekira XR™	<input type="checkbox"/> Vosevi®	<input type="checkbox"/> Zepatier®
<input type="checkbox"/> peginterferon alfa	<input type="checkbox"/> ribavirin	<input type="checkbox"/> Mavyret™	
<input type="checkbox"/> sofosbuvir (400mg)/ velpatasvir (100mg)	<input type="checkbox"/> ledipasvir (90mg)/ sofosbuvir (400mg)		

- **Treatment is being prescribed by (check box below that applies):**

<input type="checkbox"/> Gastroenterologist	<input type="checkbox"/> Hepatologist	<input type="checkbox"/> ID Specialist
<input type="checkbox"/> Transplant Specialist	<input type="checkbox"/> Other:	

- **Patient is:** treatment naïve relapse treatment experienced
- **Patient tried and failed PREFERRED drug (Mavyret™)**
- Patient has a diagnosis of chronic HCV genotype. 1 2 3 4 5 6 **(Labs must be included)**
- A documented viral load (HCV RNA) taken within 6 months of beginning therapy **(Include labs)**
- Is member co-infected with hepatitis B **(send labs)** Yes No

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- Is member co-infected with HIV-1? Yes No
- Has patient ever been successfully treated for chronic HCV? Yes No
- Does patient have hepatocellular carcinoma or severe cirrhosis awaiting a liver transplant? Yes No
- Does patient have compensated cirrhosis? **(Include labs)** Yes No
- Does patient has decompensated cirrhosis (which is defined as a Child-Pugh score greater than 6 [class B or C])? **(Include labs)** Yes No

For assessment of disease severity, please refer to the table below for a Fibrosis/stage score:

Metavir	Scheuer/Batts, Ludwig/Tsui Stage	Ishak, et al: Fibrosis Scoring	Knodell et al: Fibrosis Scoring
F0=no fibrosis	0=No fibrosis, normal amount of connective tissue	0=No fibrosis	0=No fibrosis
F1=portal fibrosis without septa	1=Portal/periportal fibrosis	1=Expansion of some portal areas with or without septa	1=Fibrous some portal expansion with or without septa
F2=portal fibrosis with rare septa	2=Septal fibrosis	2=Expansion of most portal areas with or without septa	2= Fibrous most portal expansion with or without septa
F3=numerous septa, not cirrhosis	3=Bridging fibrosis with architectural distortion.	3=Expansion of most portal areas with occasional portal to portal bridging	3=Bridging Fibrosis
F4=cirrhosis	4=Cirrhosis, probable cirrhosis	4=Expansion of portal areas with marked bridging (portal-portal and/or portal-central)	4=Cirrhosis
		5=Marked bridging with occasional nodules (incomplete cirrhosis)	
		6=Cirrhosis, probable or definitive	

Cirrhosis requires 2 liver assessments with Lab values & symptoms correlating with Cirrhosis. Submit a Liver assessment documenting fibrosis including one (1) of the following: **(Please Note: Contra-Indication to a liver assessment would lead to an incomplete form.)**

Liver biopsy confirming:

<input type="checkbox"/> METAVIR score	<input type="checkbox"/> Knodell fibrosis score (last #-reported separately)
<input type="checkbox"/> Ishak stage	<input type="checkbox"/> Batts-Ludwig stage

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- Transient elastography (FibroScan) score of: _____ kPa
- FibroTest (FibroSure) score of: _____ (Alcohol test **MUST** be same date of FibroTest)
- Shear wave elastography (ElastPQ) score of: _____ m/s
- Shear wave (SWE supersonic tech) score of: _____ m/s
- Shear wave (VTTQ) Siemens score of: _____ m/s
- LABS need to be submitted with this request form for the following: CBC BMP

****Use of samples to initiate therapy does not meet step edit/ preauthorization criteria.****

Previous therapies will be verified through pharmacy paid claims or submitted chart notes.

Patient Name: _____

Member Optima #: _____ Date of Birth: _____

Prescriber Name: _____

Prescriber Signature: _____ Date: _____

Office Contact Name: _____

Phone Number: _____ Fax Number: _____

DEA OR NPI #: _____

*Approved by Pharmacy and Therapeutics Committee: 10/19/2017
REVISED/UPDATED: ~~12/30/2017~~ 1/7/2019.

Optima Health Plan

Hepatitis C Therapy Patient Treatment Agreement

Prescriber Instructions: Please submit the completed agreement with the **initial prior authorization requests.**

Patient Instructions: By reading and signing this agreement, I acknowledge that I have been informed about the requirements of the treatment program and understand what is expected of me. I can refuse to sign this agreement, but treatment will not be started until and unless I sign this agreement.

Patient Information	Prescriber Information
Name: _____	Name: _____
Optima Health Member ID Number: _____	Optima Provider ID Number or NPI: _____
Date of Birth: _____	Office Contact Name: _____
Hepatitis C Medication Regimen: _____ _____	Telephone Number: _____ Fax Number: _____
1. I have been told how to take my hepatitis-C medicines. I understand how to take them. I am aware of possible side effects. I understand why it is important to finish all the therapy.	
2. I will take my hepatitis-C medicines like my doctor said. I understand that missing doses of medicine may cause the treatment to fail.	
3. I understand that if I miss more than 3 doses in one month, OPTIMA may no longer pay for my hepatitis-C medicines.	
4. I will tell my doctor and pharmacist the medicines I take. I understand there may be some medicines I cannot take with my hepatitis-C medicines.	
5. I understand that OPTIMA may only pay for hepatitis-C medicines for a certain number weeks over my lifetime.	
6. I understand that past use of certain hepatitis-C medicines may keep me from using medicines like them again.	
7. I am not currently using IV drugs or abusing alcohol.	
8. I will not use IV drugs or abuse alcohol (which could seriously damage my liver) while on treatment or after completion of treatment.	
9. I am (OR my female partner is) not pregnant.	
10. I am (OR my female partner is) not planning on getting pregnant while I am on my hepatitis-C medicines and for at least 6 months after I finish them.	
11. I (OR my female partner) will use two forms of non-hormonal birth control while I am taking my hepatitis-C medicines and for at least 6 months after I finish taking them.	
12. I (OR my female partner) will have monthly pregnancy testing while I am taking my hepatitis-C medicines.	

I have read the above statements and understand the agreement.

Patient Signature: _____ **Date:** _____

Physician Signature: _____ **Date:** _____