

**HEPATITIS C AGENTS  
PRIOR AUTHORIZATION FORM**  
(form effective 1/5/2026)



Fax to PerformRx<sup>SM</sup> at **1-855-851-4058**, or to speak to a representative call **1-888-674-8720**.

**PRIOR AUTHORIZATION REQUEST INFORMATION**

Office contact name/phone:	
LTC facility contact/phone:	Total # pages:

**BENEFICIARY INFORMATION**

Beneficiary name:	Beneficiary ID #:	DOB:
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**PRESCRIBER INFORMATION**

Prescriber name:		
Specialty:	NPI:	State license #:
Street address:		
City/state/zip:		
Phone:	Fax:	

**CLINICAL INFORMATION**

Requested drug #1:	
Directions:	
Qty:	<input type="checkbox"/> 8 weeks <input type="checkbox"/> 12 weeks <input type="checkbox"/> 16 weeks <input type="checkbox"/> Other:
Requested drug #2:	
Directions:	
Qty:	<input type="checkbox"/> 8 weeks <input type="checkbox"/> 12 weeks <input type="checkbox"/> 16 weeks <input type="checkbox"/> Other:
Is the beneficiary currently being treated with the requested drug(s)? <input type="checkbox"/> Yes – Therapy start date: _____ <input type="checkbox"/> No	

**SUBMIT DOCUMENTATION from the medical record for all items below.**

**For requests for NON-PREFERRED Hepatitis C Agents direct-acting antivirals (DAAs):**

- Documentation that the beneficiary tried and failed or has a contraindication or intolerance to the preferred Hepatitis C Agents. (See the Preferred Drug List for the list of preferred Hepatitis C Agents at: <https://papdl.com/preferred-drug-list>.)**  
List medications tried: \_\_\_\_\_
- Cirrhosis assessment documented by a recent noninvasive test and date of testing.**
- Genotype if one of the following (check the appropriate box for the beneficiary):**  
 The beneficiary is prescribed a non-pangenotypic regimen.  
 The beneficiary is hepatitis C sofosbuvir-based, sofosbuvir-velpatasvir-voxilaprevir, or sofosbuvir plus glecaprevir-pibrentasvir treatment-experienced.  
 The beneficiary has decompensated cirrhosis and is prescribed ledipasvir-sofosbuvir.  
 The beneficiary is treatment-naïve (with cirrhosis) and prescribed sofosbuvir-velpatasvir.
- RAS (resistance-associated substitutions) testing and date of testing if one of the following (check the appropriate box for the beneficiary):**  
 The beneficiary is genotype 1a and prescribed elbasvir-grazoprevir.  
 The beneficiary is genotype 1a, treatment-experienced, and prescribed ledipasvir-sofosbuvir.  
 The beneficiary is genotype 3, treatment-naïve (with cirrhosis) or treatment-experienced (without cirrhosis) and prescribed 12 weeks of sofosbuvir-velpatasvir.

**For requests for THERAPEUTIC DUPLICATION of Hepatitis C Agents direct-acting antivirals (DAAs):**  
For a beneficiary taking more than 1 Hepatitis C Agents DAA product concomitantly:  
 The beneficiary has a medical reason for concomitant use of the requested products that is supported by peer-reviewed medical literature or national treatment guidelines.

**For requests for ALL OTHER NON-PREFERRED Hepatitis C Agents (e.g., Pegasys):** Diagnosis: \_\_\_\_\_  
 The beneficiary has a history of therapeutic failure of or a contraindication or an intolerance to first line therapies.  
List therapies tried: \_\_\_\_\_

**ATTESTATION from the prescriber for one of the items below.**

**Check the appropriate box for the beneficiary.**  
 The beneficiary is hepatitis C treatment naïve.  
 The beneficiary has been treated for hepatitis C with the following treatment regimen: \_\_\_\_\_

**PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION**

Prescriber signature:	Date:
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