MASSACHUSETTS STANDARD FORM FOR HEPATITIS C MEDICATION PRIOR AUTHORIZATION REQUESTS

*Some plans might not accept this form for Medicare or Medicaid requests.

A. Destination		
Health Plan or Prescription Plan Name: Harvard Pilgrim Health Car	e, Attn: Pharmacy Utilization Management Dept	
Health Plan Phone: 1-800-708-4414 Health Plan Fax: 1-617-673-0	Online Prior Authorization: https://point32health.promptpa.com	
B. Patient Information		
Patient Name: DOB:	Gender: Male Female Other:	
Member ID #:		
C. Prescriber Information		
Prescribing Clinician:	Phone #:	
Specialty:	Secure Fax #:	
NPI #:	DEA #:	
Prescriber Point of Contact Name (POC) (if different than prescriber):		
POC Phone #:	POC Secure Fax #:	
POC Email (not required):		
Prescribing Clinician or Authorized Representative Signature:		
Date:		
D. Medication Information		
Check if Expedited Review/Urgent Request: [In checking this box, I attest to the fact that this request meets the definition and criteria for expedited review and is an urgent request.)		
□ Daklinza □ Epclusa □ Harvoni □ Olysio □ Ribavirin Generic □ Ribavirin Branded		
☐ Sovaldi ☐ Technivie ☐ Viekira Pak ☐ Viekira XR ☐ Zepatier ☐ Vosevi ☐ Mavyret ☐ Other		
Requested Duration of Treatment: weeks		
Type of Therapy: Initial Continuation — weeks remaining:		
Anticipated or actual start date:		
Is the medication prescribed by, or in consultation with, a gastroenterologist, infectious disease specialist, or hepatologist?		
For Zepatier only: Has there been confirmation that the patient does not have a genotype 1a with a baseline NS5A polymorphism?		
Yes No Unknown		
For Ribavirin only: Does the patient require a dosage form other than generic ribavirin 200 mg capsules or tablets? Yes No If yes, please specify the following:		
Dosage form requested:		
Clinical reason for use:		
Are any of the following statements true?		
☐ Patient is pregnant or plans to become pregnant within 6 months of completing treatment		
Patient is male with a female partner who is pregnant or plans to become pregnant within 6 months of completing treatment		
Patient has contraindications or intolerance to Ribavirin		

(continued on next page)

E. Patient Clinical Information			
*Please refer to plan-specific criteria for details related to required information.			
Diagnosis: B18.2 Hepatitis C (chronic) Other:			
HCV Genotype: ☐ 1 ☐ 1a ☐ 1b ☐ 2 [3	Stage of Hepatic Fibrosis: F0 F1 F2 F3 F4	
		If F4: ☐ Compensated ☐ Decompensated	
Check all methods of assessment that apply	and include result:		
Method		Result	
Liver biopsy		See above	
☐ Transient elastography (FibroScan)		kPa	
☐ Shear wave elastography		kPa	
☐ MRE		kPa	
☐ FibroSure (FibroTest)			
☐ Echosens Fibrometer			
Fibrospect			
☐ APRI			
□ Fib-4			
☐ Hepascore ☐ Other:			
Does the patient have HIV coinfection? Yes No Unknown			
Is the patient status post liver transplant? Yes No			
	15−29	se specify.)	
HCV RNA levels:	W1/ 1		
Baseline (most recent):	IU/mL Date of	of lab work: IU/mL	
week of treatment (if continuation request).			
Previous Treatments Has the patient been previously treated for Hepatitis C and failed treatment? ☐ Yes ☐ No			
	atitis C and falled treatment? L	j řes 🔲 no	
Adverse Reaction? Yes No			
Drug Name	Date of treatment (MM/YY)	Response to treatment	
		Relapsed	
		Partial response	
		☐ Null response (<2 log reduction in HCV RNA at Week 12) ☐ Did not complete	
		Briefly describe details:	
		☐ Relapsed☐ Partial response	
		Null response (<2 log reduction in HCV RNA at Week 12)	
		☐ Did not complete	
		☐ Briefly describe details:	
		Relapsed	
		☐ Partial response	
		☐ Null response (<2 log reduction in HCV RNA at Week 12)	
		☐ Did not complete	
		☐ Briefly describe details:	
Additional information pertinent to this request:			

Providers should consult the health plan's coverage policies, member benefits, and medical necessity guidelines to complete this form.

Providers may attach any additional data relevant to medical necessity criteria.