

State of Oklahoma **Oklahoma Health Care Authority**

Mavyret™ (Glecaprevir/Pibrentasvir) Initiation Prior Authorization Form

Member Name:		Date of Birth:	Member ID#:	
Pharmacy NPI:		Date of Birth: Pharmacy Phone:	Pharmacy Fax:	
Pharmacy Name:		Pharmacist Nam	Pharmacist Name:	
F	Prescriber NPI:	Prescriber Name:	Specialty:	
		Prescriber Fax:	Drug Name:	
NDC: Start Date:				
	Clinical Information			
2.	METAVIR Equivalent Date Fibrosis Stage	uding subtype if applicable): l nt Fibrosis Stage: Testing Type: Determined:		
3.	Pre-Treatment Viral	Load: Date Determined:		
4.	C)? Yes No_	decompensated hepatic disease or moderate-to-	severe nepatic impairment (Child-Pugh B or	
5.	Is the member curre	ntly on hospice or does the member have a limited by treating HCV? Yes No	ed life expectancy (less than 12 months) that	
6.	. Has the member been evaluated by a gastroenterologist, infectious disease specialist, or a transplant specialist within the past 3 months? Yes No			
	7. If yes, please include name of specialist recommending hepatitis C treatment:			
	. Has the member been previously treated for hepatitis C? Yes No Did the member's prior treatment regimen contain an NS5A inhibitor (e.g., daclatasvir, elbasvir, ledipasvir, om-			
9.	bitasvir, pibrentasvir, velpatasvir)? Yes No			
10	10. Did the member's prior treatment regimen contain an NS3/4A protease inhibitor (e.g., boceprevir, glecaprevir, grazo			
previr, paritaprevir, simeprevir, telaprevir, voxilaprevir)? Yes No				
11. Please indicate previous treatment regimen and reason for failure (relapser, null-responder, partial responder):				
	 Mavyret™ 10 Mavyret™ 10 Mavyret™ 10 Other: 	nested regimen below: 00mg/40mg daily x 56 days (8 weeks) 00mg/40mg daily x 84 days (12 weeks) 00mg/40mg daily x 112 days (16 weeks)		
 13. Has the member signed the intent to treat contract**? Yes No **Required for processing of request 14. Has the member been counseled on the harms of illicit IV drug use and alcohol use and agreed to not use illicit IV drugs or alcohol while on or after they finish hepatitis C treatment? Yes No 15. Has the member initiated immunization with the hepatitis A and B vaccines? Yes No 16. For women of childbearing potential (and male patients with female partners of childbearing potential): Patient is not pregnant (or a male with a pregnant female partner) and not planning to become pregnant during treatment Agreement that partners will use two forms of effective non-hormonal contraception during treatment. Please list non-hormonal birth control options discussed with member 17. Is the member taking any of the following medications: carbamazepine, rifampin, ethinyl estradiol containing medical 				
	tions, St. John's work vastatin doses great	t, atazanavir, darunavir, lopinavir, ritonavir, efavir ter than 10mg, or cyclosporine doses greater than ally significant issues been addressed prior to star	enz, atorvastatin, lovastatin, simvastatin, rosun 100mg per day? Yes No	
10				
		d this patient be followed by an OHCA Care Mana	-	
Members must be adherent for continued approval. Treatment gaps of therapy longer than 3 days will result in denial of payment for subsequent requests for continued therapy. Refills must be prior authorized.				
Prescriber Signature: Date:				
Has the member been counseled on appropriate use of Mavyret™ therapy? Yes No Pharmacist Signature: Date:				
Pharmacist Signature: Date: Please do not send in chart notes. Failure to complete this form in full will result in processing delays. By signature, the prescriber or pharmacist confirms the above information is accurate.				

PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

University of Oklahoma College of Pharmacy **Pharmacy Management Consultants** Product Based Prior Authorization Unit Fax: 1-800-224-4014

Phone: 1-800-522-0114 Option 4

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