Prior Authorization Request Form for pemigatinib (Pemazyre)



JOHNS HOPKINS HEALTHCARE

7231 Parkway Drive, Suite 100, Hanover, MD 21076

FAX Completed Form and Applicable Progress Notes to: (410) 424-4037

USFHP Pharmacy Prior Authorization Form

To be completed by Requesting provider			
Drug Name:	Strength:		
Dosage/Frequency (SIG):	Duration of Therapy:		

Questions? Contact the Pharmacy Dept at: (888) 819-1043, option 4

Clinical Documentation must accompany form in order for a determination to be made.

Step	Please complete patient and physician information (please print):				
1	Patient Name: Ph	ress: Address: nsor ID # Phone #:			
	Address:				
	Sponsor ID #				
	Date of Birth:				
Step	se complete the clinical assessment:				
2	1. Is the patient greater than or equal to 18 years of age?	☐ Yes	□ No		
		Proceed to question 2	STOP		
			Coverage not approved		
	2. Is the requested medication prescribed by or in consultation	☐ Yes	□ No		
	with a hematologist or oncologist?	Proceed to question 3	STOP		
			Coverage not approved		
	3. What is the indication or diagnosis?	☐ Pathologically confirmed unresectable or advanced/metastatic cholangiocarcinoma with a fibroblast growth factor receptor 2 (FGFR2) fusion or other rearrangement – Proceed to question 4			
		☐ Relapsed or refractory myeloid/lymphoid neoplasms (MLNs) with FGFR1 rearrangement – Proceed to question 7			
		☐ Other- Proceed to question 5			
	4. Has the diagnosis been detected by an FDA-approved test?	☐ Yes	□ No		
		Proceed to question 7	STOP		
			Coverage not approved		
	5. Please provide the indication or diagnosis.				
		Proceed to question 6			

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	6. Is the diagnosis cited in the National Comprehensive Cancer Network (NCCN) guidelines as a category 1, 2A, or 2B recommendation?	☐ Yes Proceed to question 7	□ No STOP Coverage not approved
,	7. Will the patient be monitored for ophthalmologic disorders including pre-treatment screening for retinal disorders?	☐ Yes Proceed to question 8	□ No STOP Coverage not approved
	8. Will the patient be monitored for hyperphosphatemia?	☐ Yes Proceed to question 9	☐ No STOP Coverage not approved
	9. Is the patient of childbearing potential?	☐ Yes Proceed to question 10	☐ No Sign and date below
	10. What is the patient's gender?	☐ Male — Proceed to question☐ Female — Proceed to ques	
,	11. Will the patient use effective contraception during treatment and for at least 1 week after the cessation of therapy?	☐ Yes Sign and date below	□ No STOP Coverage not approved
,	12. Will the patient use effective contraception during treatment and for at least 1 week after the cessation of therapy?	☐ Yes Proceed to question 13	□ No STOP Coverage not approved
	13. Is the patient pregnant?	☐ Yes STOP Coverage not approved	□ No Proceed to question 14
,	14. Has it been confirmed that the patient is not pregnant by (-) HCG?	☐ Yes Proceed to question 15	□ No STOP Coverage not approved
	15. Will the patient not breastfeed during treatment and for at least 1 week after the cessation of treatment?	☐ Yes Sign and date below	□ No STOP Coverage not approved
Step 3	I certify the above is true to the best of my knowledge Please sign and date:	e.	
	Prescriber Signature	Date	[05 April 2023]
For Inte	ernal Use Only		
Approved:		Duration of Approval:month(s)	
Denied:		Authorized By:	
☐ Incor	nplete/Other:	PA#:	

Date Decision Rendered:

Date Faxed to MD: