## TRICARE Prior Authorization Request Form for avapritinib (Ayvakit)



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:		
	0.0.0.0		
Dosage/Frequency (SIG):	Duration of Therapy:		
3 1 7 7	17		

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	Dia				
<b>1</b>	Please complete patient and physician information (please print):  Patient Name:  Physician Name:				
•	Patient Name: Physician Name: Address: Address:				
	Sponsor ID #:		Phone #:		
	Date of Birth: Secure Fax #:				
Step 2	Please complete the clinical assessment:				
	1. Is the patient GREATER THAN or EQUAL to 18 years of	☐ Yes	□ No		
		age?	Proceed to question 2	STOP	
				Coverage not approved	
	2.	2. Is the requested medication prescribed by or in consultation with a hematologist/oncologist?	☐ Yes	□ No	
			Proceed to question 3	STOP	
				Coverage not approved	
	3.	Does the patient have pathologically confirmed unresectable or metastatic gastrointestinal stromal tumor	☐ Yes	□ No	
	(GIST) harboring a platelet-derived growth factor receptor alpha (PDGFRA) exon 18 mutation with or without the D842V mutation?	Proceed to question 8	Proceed to question 4		
	4.		☐ Yes	□ No	
		(includes patients with aggressive systemic mastocytosis, systemic mastocytosis with an associated hematologic neoplasm, and mast cell leukemia)?	Proceed to question 8	Proceed to question <b>5</b>	
	5. Does the patient have indolent systemic mastocytosis (ISM) with a platelet count GREATER THAN OR EQUAL TO 50 x 10^9/L?	□ Yes	□ No		
		Proceed to question 8	Proceed to question 6		
	6.	Please provide the indication or diagnosis.			
			Proceed to question 7		

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	Cancer Network (NCCN) guidelines as a category 1, 2A, or 2B recommendation?	☐ Yes Proceed to question 8	□ No STOP Coverage not approved
	8. Will the provider monitor for intracranial bleeding and other central nervous system (CNS) adverse effects?	☐ Yes Proceed to question 9	□ No STOP Coverage not approved
	9. What is the patient's gender?	☐ Male — Proceed to question 10 ☐ Female — Proceed to question 11	
	10. Will the patient use effective contraception during treatment and for at least 6 weeks after the cessation of therapy?	☐ Yes Sign and date below	□ No STOP Coverage not approved
	11. Is the patient of childbearing potential?	☐ Yes Proceed to question 12	□ No Sign and date below
	12. Will the patient use effective contraception during treatment and for at least 6 weeks 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 <b>15</b>	□ No STOP Coverage not approved
	15. Will the patient breastfeed during treatment and for at least 2 weeks after the cessation of treatment?	☐ Yes STOP Coverage not approved	□ No Sign and date below
tep 3	I certify the above is true to the best of my knowledge	Please sign and date:	
-	Prescriber Signature	Date	
			[03 January 2024]
or Inte	ernal Use Only		
] Appr	roved:	Duration of Approval:	month(s)
] Deni	ed:	Authorized By:	
] Incor	mplete/Other:	PA#:	
ate Fa	axed to MD:	Date Decision Rendered:	