

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.

	ithorization does not expire.			
Step	Please complete patient and physician information	on (please print):		
1	Patient Name:	Physician Name:		
	Address:	Address:		
	Sponsor ID #	Phone #:		
	Date of Birth:	Secure Fax #:		
Step	Please complete the clinical assessment:			
2	1. Will Tafinlar be used in combination with Mekinist (trametinib)?	☐ Yes Proceed to question 2	□ No Proceed to question 3	
	2. For which indication is Tafinlar being prescribed?	Melanoma - Proceed	Melanoma - Proceed to question 3	
		Metastatic Non-small to question 7	Metastatic Non-small Cell Lung cancer – Proceed to question 7	
		cancer without satisfactor	Locally advanced or metastatic anaplastic thyroid cancer without satisfactory locoregional treatment options - Proceed to question 7	
		Solid tumor, unresecta progression following pric satisfactory alternative tre to question 5		
		Low-grade glioma (LG therapy - Proceed to que		
		□ Other - Proceed to qu	estion 9	
	3. Does the patient have unresectable or metastatic melanoma?	□ Yes	□ No	
		Proceed to question 4	Proceed to question 9	
	4. Does the patient have a BRAF-V600E or BRAF-V600K mutation as detected by an FDA-approved test?	K 🗆 Yes	□ No	
		Proceed to question 8	Proceed to question 9	
	5. Is the patient greater than or equal to 1 year of age?	Yes	□ No	
		Proceed to question 7	Proceed to question 9	

Continue on next page

TRICARE Prior Authorization Request Form for dabrafenib **(Tafinlar)**

6. How old is the patient?	□ Less than 1 year of age - Proceed to question 9	
	□ 1 year of age or older by age - Proceed to question 7	
	□ Greater than 18 years o question 9	f age - Proceed to
7. Does the patient have a BRAF-V600E mutation as detected by an FDA-approved test (if one is available for this indication)?	□ Yes	🗆 No
	Proceed to question 8	Proceed to question 9
 Is the patient taking encorafenib (Braftovi), binimetinib (Mektovi), vemurafenib (Zelboraf), or cobimetinib (Cotellic)? 	□ Yes	🗆 No
	STOP	Sign and date below
	Coverage not approved	
9. Please provide the diagnosis.		
	Proceed to question 10	
10. Is the diagnosis cited in the National Comprehensive Cancer Network (NCCN) guidelines as a category 1, 2A, or 2B recommendation?	□ Yes	🗆 No
	Sign and date below	STOP
		Coverage not approved

Step I certify the above is true to the best of my knowledge. Please sign and date:
3

	D 1 L	
Prescriber Sidnature	Date	
	Duto	

[03 January 2024]

For Internal Use Only				
Approved:	Duration of Approval:month(s)			
Denied:	Authorized By:			
Incomplete/Other:	PA#:			
Date Faxed to MD:	Date Decision Rendered:			