## USFHP Prior Authorization Request Form for trametinib (**Mekinist**)

To be completed and signed by the prescriber. To be used only for prescriptions which are to be filled through the Department of Defense (DoD) US Family Health Plan Pharmacy Program. US Family Health Plan is a TRICARE contractor for DoD.

The completed form may be faxed to 855-273-5735

**OR** 

The patient may attach the completed form to the prescription and **mail** it to:

Attn: Pharmacy, 77 Warren St, Brighton, MA 02135

QUESTIONS? Call 1-877-880-7007

https://www.usfamilyhealth.org/for-providers/pharmacy-information/

Step	Please complete patient and physician inform	nation (please print):			
1	Patient Name:	Physician Name:			
	Address:	Address:			
	Sponsor ID #	Phone #:			
	Date of Birth:	Secure Fax #:			
Step 2	Please complete the clinical assessment:				
	Is the requested medication prescribed by or in consultation with a hematologist/oncologist?	□ Yes	□ No		
		Proceed to question 2	STOP		
			Coverage not approved		
	2. Will Mekinist be used in combination with Tafinlar (dabrafenib)?	. D Yes	□ No		
		Proceed to question 3	Proceed to question 4		
	3. For which indication is Mekinist being prescribed?	?	☐ Melanoma – Proceed to <b>6</b>		
			☐ Adjuvant treatment of patients of melanoma with involvement of lymph node(s), following complete resection - Proceed to 6		
		☐ Metastatic Non-small question <b>7</b>	☐ Metastatic Non-small Cell Lung cancer – Proceed to question <b>7</b>		
		can cer without satisfacto	☐ Locally advanced or metastatic anaplastic thyroid cancer without satisfactory locoregional treatment options - Proceed to question 7		
			☐ Low-grade glioma (LGG) requiring systemic therapy - Proceed to question 7		
		progression following	☐ Solid tumor, unresectable or metastatic, with progression following prior treatment and no satisfactory alternative treatment options - Proceed to question 7		
		☐ Other - Proceed to qu	☐ Other - Proceed to question 9		
	4. Has the patient received prior BRAF-inhibitor therapy, for example, with Tafinlar or Zelboraf?	apy, 🗆 Yes	□ No		
		Proceed to question §	Proceed to question 5		

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	5. Does the patient have unresectable or metastatic melanoma?	□ Yes	□ No	
	mounoma:	Proceed to question 6	Proceed to question 9	
	6. Does the patient have a BRAF-V600E or BRAF-V600K mutation as detected by an FDA-approved test?	□ Yes	□ No	
	matation as assessed by an i BA approved test.	Proceed to question 8	Proceed to question 9	
	7. Does the patient have a BRAF-V600E mutation as detected by an FDA-approved test (if one is available	☐ Yes	□ No	
	for this indication)?	Proceed to question 8	Proceed to question 9	
	8. Is the patient taking encorafenib (Braftovi), binimetinib (Mektovi), vemurafenib (Zelboraf), or cobimetinib (Cotellic)?	□ Yes	□ No	
		STOP	Sign and date below	
		Coverage not approved		
	<ol> <li>The diagnosis IS NOT listed above but IS cited in the National Comprehensive Cancer Network (NCCN) guidelines as a category 1, 2A, or 2B recommendation.</li> </ol>			
	To facilitate approval, please list the diagnosis, guideline version, and page number:			
		Sign and date below		
Step 3	I certify the above is true to the best of my knowledge. Please sign and date:			
	Prescriber Signature	 Date		
	1 1000 Dor Orginataro	Date	[08 January 2025]	