

US Family Health Plan
 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**

Prior Authorization does not expire.

Step 1 Please complete patient and physician information (please print):

Patient Name: _____ Address: _____ Sponsor ID #: _____ Date of Birth: _____	Physician Name: _____ Address: _____ Phone #: _____ Secure Fax #: _____
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Step 2 Please complete the clinical assessment:

1. Will Mekinist be used in combination with Tafinlar (dabrafenib)?	<input type="checkbox"/> Yes Proceed to question 2	<input type="checkbox"/> No Proceed to question 3
2. For which indication is Mekinist being prescribed?	<input type="checkbox"/> Melanoma - Proceed to 4 <input type="checkbox"/> Metastatic Non-small Cell Lung cancer – Proceed to question 8 <input type="checkbox"/> Locally advanced or metastatic anaplastic thyroid cancer without satisfactory locoregional treatment options - Proceed to question 8 <input type="checkbox"/> Low-grade glioma (LGG) requiring systemic therapy - Proceed to question 7 <input type="checkbox"/> Solid tumor, unresectable or metastatic, with progression following prior treatment and no satisfactory alternative treatment options - Proceed to question 6 <input type="checkbox"/> Other - Proceed to question 10	
3. Has the patient received prior BRAF-inhibitor therapy, for example, with Tafinlar or Zelboraf?	<input type="checkbox"/> Yes Proceed to question 10	<input type="checkbox"/> No Proceed to question 4
4. Does the patient have unresectable or metastatic melanoma?	<input type="checkbox"/> Yes Proceed to question 5	<input type="checkbox"/> No Proceed to question 10
5. Does the patient have a BRAF-V600E or BRAF-V600K mutation as detected by an FDA-approved test?	<input type="checkbox"/> Yes Proceed to question 9	<input type="checkbox"/> No Proceed to question 10

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6. Is the patient greater than or equal to 1 year of age?	<input type="checkbox"/> Yes Proceed to question 8	<input type="checkbox"/> No Proceed to question 10
7. How old is the patient?	<input type="checkbox"/> Less than 1 year of age - Proceed to question 10 <input type="checkbox"/> 1 year of age or older but less than 18 years of age - Proceed to question 8 <input type="checkbox"/> Greater than 18 years of age - Proceed to question 10	
8. Does the patient have a BRAF-V600E mutation as detected by an FDA-approved test (if one is available for this indication)?	<input type="checkbox"/> Yes Proceed to question 9	<input type="checkbox"/> No Proceed to question 10
9. Is the patient taking encorafenib (Braftovi), binimetinib (Mektovi), vemurafenib (Zelboraf), or cobimetinib (Cotellic)?	<input type="checkbox"/> Yes STOP Coverage not approved	<input type="checkbox"/> No Sign and date below
10. Please provide the diagnosis.	<hr style="width: 80%; margin: 0 auto;"/> Proceed to question 11	
11. Is the diagnosis cited in the National Comprehensive Cancer Network (NCCN) guidelines as a category 1, 2A, or 2B recommendation?	<input type="checkbox"/> Yes Sign and date below	<input type="checkbox"/> No STOP Coverage not approved

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

_____ Prescriber Signature

_____ Date