

Priority Health Medicare prior authorization form

Fax completed form to: 877.974.4411 toll free, or 616.942.8206 Medicare Part D This form applies to: ☐ Standard request This request is: **Expedited request** Your request will be expedited if you haven't gotten the prescription and Priority Health Medicare determines, or your prescriber tells us, that your life or health may be at risk by waiting. Braftovi[™] + Mektovi[®] (encorafenib / binimetinib) Member First Name: Last Name: DOB: _____ Gender: _ Primary Care Physician: Prov. Phone: Prov. Fax: Requesting Provider: Provider Address: Provider NPI: _____ Contact Name: Provider Signature: **Product Information** □ New request □ Continuation request Start date (or date of next dose): Drug product: Braftovi 50 mg capsule Date of last dose (if applicable): Braftovi 75 mg capsule ☐ Mektovi 15 mg capsule Dosing frequency: Prior authorization criteria The following requirements need to be met before this drug is covered by Priority Health Medicare. These requirements have been approved by the Centers for Medicare and Medicaid Services (CMS), but you may ask us for an exception if you believe one or more of these requirements should be waived. For this drug to be covered, the patient must meet the following criteria: 1. Must be used for a medically-accepted indication* 2. Must have documentation of BRAF weather mutation status as detected by an FDA-approved test 3. Patient must have an Eastern Cooperative Oncology Group (ECOG) performance status of 0 to 2

Medically-accepted indication*

This drug is only covered under Medicare Part D when it is used for a medically accepted indication. A medically accepted indication for a drug or biologic used in an anti-cancer chemotherapeutic regimen is a use that is either.

- approved by the Food and Drug Administration. (That is, the Food and Drug Administration has approved the drug for the diagnosis or condition for which it is being prescribed.)
- supported by one of the following references (known as compendia): National Comprehensive Cancer Network (NCCN) Drugs and Biologics Compendium, Micromedex DrugDex, American Hospital Formulary Service-Drug Information, Clinical Pharmacology, or Lexi-Drugs
- or supported in peer-reviewed medical literature appearing in regular editions of approved publications



Additional information
Note: When criteria are met, coverage duration is 1 year.
Priority Health Precertification Documentation
A. What condition is this drug being requested for? Unresectable or metastatic melanoma with BRAF V600K or V600E mutation Other – the patient's condition is:
B. Has the BRAF V600 mutation been detected by an FDA-approved test (documentation required)? Yes. No. Are you requesting an exception to the criteria? Yes. Rationale for exception: No
C. Is Braftovi being used in combination with Mektovi? Yes. No. Are you requesting an exception to the criteria? Yes. Rationale for exception:
 No D. What is the patient's ECOG performance status? □ 0: Fully active, able to carry on all pre-disease performance without restriction □ 1: Restricted in physically strenuous activity but ambulatory and able to carry out work of a light or sedentary nature (e.g. light house work, office work) □ 2: Ambulatory and capable of all self-care but unable to carry out any work activities; up and about more than 50% of waking hours □ 3: Capable of only limited self-care, confined to bed or chair more than 50% of waking hours □ 4: Completely disabled; cannot carry on any self-care; totally confined to bed or chair
Priority Health Medicare Exception Request (exceptions to the above criteria) Do you believe one or more of the prior authorization requirements should be waived? Yes No If yes, you must provide a statement explaining the medical reason why the exception should be approved.
Would Braftovi + Mektovi likely be the most effective option for this patient? ☐ Yes ☐ No If yes, please explain why: ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐
If the patient is currently using Braftovi + Mektovi, would changing the patient's current regimen likely result in adverse effects for the patient? Yes No If yes, please explain: