BlueCross BlueShield

RETEVMO

Send completed form to: Service Benefit Plan Prior Approval P.O. Box 52080 MC 139 Phoenix, AZ 85072-2080 Attn. Clinical Services Fax: 1-877-378-4727

Federal Employee Program. **PRIOR APPROVAL REQUEST** Additional information is required to process your claim for prescription drugs. Please complete the patient portion, and have the prescribing physician complete the

physician portion and submit this completed			F		Fax:	1-877-378-4727	
Patient Information (required)			Provider Information (required)				
Date:			Provider Name:				
Patient Name:			Specialty:		NPI:		
Date of Birth:	Date of Birth:Sex:MaleFemale		Office Phone: O		Office Fax:	Office Fax:	
Street Address:			Office Street Addres	ss:			
City:	State:	Zip:	City:	Sta	te:	Zip:	
Patient ID: R			Physician Signature	:		1	
		PHYSICIAN	COMPLETES				
Retevmo (selpercatinib)							

**Check www.fepblue.org/formulary to confirm which medication is part of the patient's benefit

NOTE: Form must be completed in its entirety for processing

Is this request for brand or generic? Brand Generic

Will the patient need more than 320 milligrams per day? □Yes* □No

*If YES, please specify the requested milligrams per day: _____ mg per day

1. Has the patient been on Retevmo continuously for the last 6 months excluding samples? Please select answer below:

NO – this is **INITIATION** of therapy, please answer the following questions:

- a. What is the patient's diagnosis?
 - □Locally advanced non-small cell lung cancer (NSCLC) <u>OR</u> □Metastatic non-small cell lung cancer (NSCLC) i. Is the patient's diagnosis RET fusion-positive, as detected by an FDA-approved test? □Yes □No
 - Advanced medullary thyroid cancer (MTC) <u>OR</u> Metastatic medullary thyroid cancer (MTC)
 - i. Does the patient have RET-positive mutation, as detected by an FDA-approved test? \Box Yes \Box No ii. Does the patient require systemic therapy? \Box Yes \Box No

Advanced thyroid cancer **OR** Metastatic thyroid cancer

- i. Is the patient's diagnosis RET fusion-positive, as detected by an FDA-approved test? Yes No
- ii. Is the patient radioactive iodine-refractory, if radioactive iodine is appropriate? *Please select answer below:*
 - Yes No Radioactive iodine is not appropriate
- iii. Does the patient require systemic therapy? **D**Yes **D**No

Locally advanced solid tumors **OR OR O**Metastatic solid tumors

i. Is the patient's diagnosis RET fusion-positive, as detected by an FDA-approved test? Ues No

ii. Has the patient's disease progressed on or following prior systemic treatment? Yes No*

*If NO, does the patient have satisfactory alternative treatment options? \Box Yes \Box No

Other diagnosis (*please specify*):

b. Will any pre-existing hypocalcemia, hypokalemia, or hypomagnesemia be corrected prior to staring Retevmo? Tyes No

□ YES – this is a PA renewal for CONTINUATION of therapy, please answer the following questions:

a. What is the patient's diagnosis?

Locally advanced non-small cell lung cancer (NSCLC)	<u>OR</u>	□ Metastatic non-small cell lung cancer (NSCLC)
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Advanced medullary thyroid cancer (MTC) **OR** Imetastatic medullary thyroid cancer (MTC)

 $\Box Advanced thyroid cancer \quad \underline{OR} \quad \Box Metastatic thyroid cancer$

□Locally advanced solid tumors <u>OR</u> □Metastatic solid tumors

Other diagnosis (please specify): _

b. Has the patient experienced disease progression or unacceptable toxicity while on Retevmo? Yes No

PLEASE PROCEED TO PAGE 2 FOR ADDITIONAL QUESTIONS

The information provided on this form will be used to determine the provision of healthcare benefits under a U.S. federal government program, and any falsification of records may subject the provider to prosecution, either civilly or criminally, under the False Claim Acts, the False Statements Act, the mail or wire fraud statutes, or other federal or state laws prohibiting such falsification. **Prescriber Certification:** I certify all information provided on this form to be true and correct to the best of my knowledge and belief. I understand that the insurer may request a medical record if the information provided herein is not sufficient to make a benefit determination or requires clarification and l agree to provide any such information to the insurer. Retevmo – FEP MD Fax Form Revised 06/28/2024



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PAGE 2 - PHYSICIAN COMPLETES

Patient Name: ____

_ DOB: __

___ Patient ID: R ____

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2. Does the prescriber agree to monitor AST, ALT, and blood pressure? Yes No

3. Does the prescriber agree to monitor for QT interval prolongation? \Box Yes \Box No

- 4. FEMALE Patient: Is the patient of reproductive potential? □Yes* □No
 *If YES, will the patient be advised to use effective contraception during treatment with Retevmo and for one week after the last dose? □Yes □No
- 5. MALE Patient: Does the patient have a female partner of reproductive potential? □Yes* □No
 *If YES, will the patient be advised to use effective contraception during treatment with Retevmo and for one week after the last dose? □Yes □No

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BlueShield. RETEVMO Federal Employee Program. PRIOR APPROVAL REQUEST

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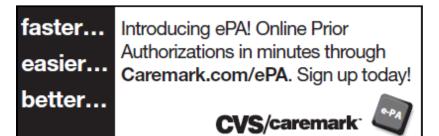
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Message:

Attached is a Prior Authorization request form.

For your convenience, there are 3 ways to complete a Prior Authorization request:

Electronically Online (ePA) Results in 2-3 minutes FASTEST AND EASIEST	Now you can get responses to drug prior authorization requests securely online. Online submissions may receive instant responses and do not require faxing or phone calls. Requests can be made 24 hours a day, 7 days a week. For more information on electronic prior authorization (ePA) and to register, go to Caremark.com/ePA.
Phone (4-5 minutes for response)	The FEP Clinical Call Center can be reached at (877)-727-3784 between the hours of 7AM- 9PM Eastern Time. A live representative will assist with the Prior Authorization, asking for the same info contained on the attached form. Please review the form and have your answers ready for faster service. The process over the phone takes on average between 4 and 5 minutes.
Fax (3-5 days for response)	Fax the attached form to (877)-378-4727 Requests sent via fax will be processed and responded to within 5 business days. The form must be filled out completely, if there is any missing information the PA request cannot be processed. <u>Please only fax the completed form once as</u> <u>duplicate submissions may delay processing</u> <u>times.</u>



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