BlueCross BlueShield Federal Employee Program.

FRUZAQLA
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 form.

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

Patient Information (required)			**Provider Information (required)**		
Date:			Provider Name:		
Patient Name:			Specialty:	NPI:	
Date of Birth:	Sex: ❑Male ❑Female		Office Phone:	Office Fax:	
Street Address:			Office Street Address:		
City:	State:	Zip:	City:	State:	Zip:
Patient ID: **R**			Physician Signature:		
PHYSICIAN COMPLETES					

Fruzaqla (fruquintinib)

NOTE: Form must be completed in its **entirety** for processing

Please select strength: ❑**1 mg quantity _____ per 84 days** ❑**5 mg quantity _____ per 84 days**

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

Is this request for brand or generic? ❑Brand ❑Generic

1. Does the patient have a diagnosis of metastatic colorectal cancer (mCRC)? ❑Yes ❑No

2. **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 Fruzaqla and for 2 weeks after the last dose? ❑Yes ❑No

3. **FEMALE Patient**: Is the patient of reproductive potential? ❑Yes* ❑No
   If YES, will the patient be advised to use effective contraception during treatment with Fruzaqla and for 2 weeks after the last dose? ❑Yes ❑No

4. Has the patient been on this medication continuously for the last **6 months** excluding samples? ***Please select answer below:***

   ❑**NO** – this is **INITIATION** of therapy, please answer the following questions:
   a. Does the patient have RAS wild-type metastatic colorectal cancer? ❑Yes* ❑No
      If YES, if medically appropriate, has the patient been previously treated with anti-EGFR therapy? ***Select answer below:***
      ❑Yes ❑No, not medically appropriate
      ❑Medically appropriate however patient has not been previously treated with anti-EGFR therapy
   b. Has the patient been previously treated with fluoropyrimidine-, oxaliplatin-, and irinotecan-based chemotherapy? ❑Yes ❑No
   c. Has the patient been previously treated with an anti-VEGF therapy? ❑Yes ❑No

   ❑**YES** – this is a PA renewal for **CONTINUATION** of therapy, please answer the following question:
   a. Has the patient experienced disease progression or unacceptable toxicity while on the requested therapy? ❑Yes ❑No

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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 information 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 Prior Authorization request cannot be processed. ***Please only fax the completed form once as duplicate submissions may delay processing times.***