

## Truseltiq (infigratinib) Prior Authorization Request Form



Caterpillar Prescription Drug Benefit Phone: 877-228-7909 Fax: 800-424-7640

**Instructions:** Please fill out all applicable sections completely and legibly. Attach any additional documentation that is important for the review (e.g., chart notes or lab data, to support the authorization request). Information contained in this form is Protected Health Information under HIPAA.

	☐ URGENT		
MEMBER INFORMATION			
LAST NAME:	FIRST NAME:		
PHONE NUMBER:	DATE OF BIRTH:		
STREET ADDRESS:			
CITY:	STATE: ZIP CODE:		
PATIENT INSURANCE ID NUMBER:			
MALE FEMALE HEIGHT (IN/CM): WEIGHT (LB/KG): ALLERGIES:  IF YOU ARE NOT THE PATIENT OR THE PRESCRIBER, YOU WILL NEED TO SUBMIT A PHI DISCLOSURE AUTHORIZATION FORM WITH THIS REQUEST WHICH CAN BE FOUND AT THE FOLLOWING LINK: https://magellanrx.com/member/external/commercial/common/doc/en-us/phi disclosure authorization.pdf			
PATIENT'S AUTHORIZED REPRESENTATIVE (IF APPLICABLE):			
PRESCRIBER INFORMATION			
LAST NAME:	FIRST NAME:		
PRESCRIBER SPECIALTY:	EMAIL ADDRESS:		
NPI NUMBER:	DEA NUMBER:		
PHONE NUMBER:	FAX NUMBER:		
STREET ADDRESS:			
CITY:	STATE: ZIP CODE:		
REQUESTOR (if different than prescriber):	OFFICE CONTACT PERSON:		
MEDICATION OR MEDICAL DISPENSING INFORMATION			
MEDICATION NAME:			
DOSE/STRENGTH: FREQUENCY:	LENGTH OF QUANTITY: THERAPY/REFILLS:		
☐ NEW THERAPY ☐ RENEWAL  DURATION OF THERAPY (SPECIFIC DATES):	IF RENEWAL: DATE THERAPY INITIATED:		

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1. HAS THE PATIENT TRIED ANY OTHER	R MEDICATIONS FOR THIS CONDITION?	YES (if yes, complete below) NO	
MEDICATION/THERAPY (SPECIFY DRUG NAME AND DOSAGE):	<b>DURATION OF THERAPY</b> (SPECIFY DATES):	RESPONSE/REASON FOR FAILURE/ALLERGY:	
2. LIST DIAGNOSES:		ICD-10:	
□ Unresectable locally advanced or met □ Other diagnosis:ICD-	10		
<b>3. REQUIRED CLINICAL INFORMATION:</b> PRIOR AUTHORIZATION.	: PLEASE PROVIDE ALL RELEVANT CLINICA	AL INFORMATION TO SUPPORT A	
Clinical Information: Is this drug being prescribed to this pa trial? □ Yes □ No	tient as part of a treatment regimen spo	ecified within a sponsored clinical	
Does patient have a fibroblast growth factor receptor 2 (FGFR2) fusion or other FGFR genetic rearrangement?  □ Yes □ No Please submit documentation.			
Does patient have cancer of the gallbladder or ampulla of Vater? ☐ Yes ☐ No			
Has patient been previously treated with a cisplatin-gemcitabine-containing regimen or a gemcitabine-containing regimen, if intolerant to cisplatin therapy?   Yes  No Please submit documentation.			
Has patient been previously treated with a MEK inhibitor such as Mekinist(trametinib), Mektovi(binimetinib), Koselugo(selumetinib) or Cotellic(cobimetinib)? ☐ Yes ☐ No			
Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?			
<b>Please note:</b> Not all drugs/diagnosis are information is received.	e covered on all plans. This request may	be denied unless all required	
the Health Plan, insurer, Medical Group	n provided is true and accurate to the beson or its designees may perform a routine uracy of the information reported on this	audit and request the medical	
Prescriber Signature or Electronic I.D.	Verification:	Date:	
you are not the intended recipient, you are here	ompanying this transmission contain confidential eby notified that any disclosure, copying, distribut have received this information in error, please no	tion, or action taken in reliance on the contents	

**FAX THIS FORM TO: 800-424-7640** 

MAIL REQUESTS TO: Magellan Rx Management Prior Authorization Program

Attn: CP - 4201 P.O. Box 64811 St. Paul, MN 55164-0811

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and arrange for the return or destruction of these documents.

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