

Kisqali (ribociclib)
Prior Authorization Request Form

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

MEMBER'S LAST NAME: _____ MEMBER'S FIRST NAME: _____

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: [PRIMETHERAPEUTICS.COM/NOPP](https://www.primetherapeutics.com/nopp)

PATIENT'S AUTHORIZED REPRESENTATIVE (IF APPLICABLE): _____
AUTHORIZED REPRESENTATIVE'S PHONE NUMBER: _____

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 THERAPY/REFILLS:	QUANTITY:
<input type="checkbox"/> NEW THERAPY	<input type="checkbox"/> RENEWAL	IF RENEWAL: DATE THERAPY INITIATED:	
DURATION OF THERAPY (SPECIFIC DATES):			

Continued on next page

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1. HAS THE PATIENT TRIED ANY OTHER MEDICATIONS FOR THIS CONDITION? <input type="checkbox"/> YES (if yes, complete below) <input type="checkbox"/> NO		
MEDICATION/THERAPY (SPECIFY DRUG NAME AND DOSAGE):	DURATION OF THERAPY (SPECIFY DATES):	RESPONSE/REASON FOR FAILURE/ALLERGY:
2. LIST DIAGNOSES:		ICD-10:
<input type="checkbox"/> Breast cancer <input type="checkbox"/> Other Diagnosis _____ ICD-10 Code(s): _____		
3. REQUIRED CLINICAL INFORMATION: PLEASE PROVIDE ALL RELEVANT CLINICAL INFORMATION TO SUPPORT A PRIOR AUTHORIZATION.		
Clinical Information: Initial Request: Is Kisqali(ribociclib) going to be used in conjunction with a clinical study? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have a diagnosis of hormone receptor (HR)-positive, human epidermal growth factor receptor-2 (HER-2)-negative advanced or metastatic breast cancer? <input type="checkbox"/> Yes <input type="checkbox"/> No Please provide chart documentation. Is patient POST-menopausal? <input type="checkbox"/> Yes <input type="checkbox"/> No Is patient PRE-menopausal or PERI-menopausal? <input type="checkbox"/> Yes <input type="checkbox"/> No Is patient a male? <input type="checkbox"/> Yes <input type="checkbox"/> No Has the patient received a previous treatment with palbociclib (Ibrance®)? <input type="checkbox"/> Yes <input type="checkbox"/> No Please provide chart documentation. Has the patient received previous treatment with abemaciclib (Verzenio®)? <input type="checkbox"/> Yes <input type="checkbox"/> No Please provide chart documentation. Has the patient progressed on first-line neoadjuvant/ adjuvant endocrine therapy for metastatic disease? <input type="checkbox"/> Yes <input type="checkbox"/> No Please provide chart documentation. Has the patient received any previous systemic chemotherapy for advanced disease? <input type="checkbox"/> Yes <input type="checkbox"/> No Please provide chart documentation. Has the patient received any previous endocrine therapy for advanced disease? <input type="checkbox"/> Yes <input type="checkbox"/> No Please provide chart documentation. Has patient received endocrine therapy for advanced disease within the past one month of requesting Kisqali? <input type="checkbox"/> Yes <input type="checkbox"/> No Has patient received more than one previous systemic chemotherapy for advanced disease? <input type="checkbox"/> Yes <input type="checkbox"/> No Will the patient use the aromatase inhibitor letrozole in combination with Kisqali (ribociclib)? <input type="checkbox"/> Yes <input type="checkbox"/> No		

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- Has the patient been previously treated with fulvestrant? Yes No
- Will the patient be using fulvestrant in combination with Kisqali? Yes No
- Will the patient be using goserelin in combination with Kisqali? Yes No
- Will the patient be using tamoxifen in combination with Kisqali? Yes No
- Will the patient be using anastrozole in combination with Kisqali? Yes No

Renewal Request:

Is the patient continuing to have a positive clinical response? Yes No Please provide chart documentation.

Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note: Not all drugs/diagnosis are covered on all plans. This request may be denied unless all required information is received.

ATTESTATION: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan, insurer, Medical Group or its designees may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.

Prescriber Signature or Electronic I.D. Verification: _____ **Date:** _____

CONFIDENTIALITY NOTICE: The documents accompanying this transmission contain confidential health information that is legally privileged. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution, or action taken in reliance on the contents of these documents is strictly prohibited. If you have received this information in error, please notify the sender immediately (via return FAX) and arrange for the return or destruction of these documents.

FAX THIS FORM TO: 800-424-7640
MAIL REQUESTS TO: Prime Therapeutics Management Prior Authorization Program
Attn: CP - 4201
P.O. Box 64811
St. Paul, MN 55164-0811