

## **CDK** Inhibitors

Please provide the information below, please print your answer, attach supporting documentation, sign, date, and return to our office as soon as possible.

Please FAX responses to: (800) 869-7791. Phone: (855) 322-4082

Date of request:					
Patient Date of birth		Мо	Molina ID		
Pharmacy name	Pharmacy NPI	Telephone number		Fax number	
Prescriber	Prescriber NPI	Telephone number		Fax number	
Medication and strength		Directions for use		Qty/Days supply	
<ol> <li>Is this request for a continuation of existing therapy? Yes No If yes, is there documentation demonstrating disease stability or a positive clinical response? Yes No</li> <li>Indicate patient's diagnosis: Adjuvant therapy of early-stage (stage I-III) breast cancer (EBC) Systemic therapy of recurrent, advanced, or metastatic breast cancer</li> </ol>					
Other, specify:					
3. Indicate stage:					
<ul> <li>4. What is the patients hormone reception and HER2 status?</li> <li>Hormone receptor: Positive Negative</li> <li>HER2: Positive Negative</li> </ul>					
5. Is this being prescribed by or in consultation with an oncologist?					
<ul> <li>6. Will this medication be used in combination with other agents for the treatment of this diagnosis?</li> <li>No</li> <li>Yes, specify regimen:</li> </ul>					
<ul> <li>7. Has patient previously progressed on, or after treatment with another CDK4/6 inhibitor (e.g., ribociclib [Kisqali], abemaciclib [Verzenio])?</li> <li>No</li> <li>Yes, explain:</li> </ul>					

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Request for Adjuvant therapy of early-stage (stage I-III) breast cancer (EBC) answer the				
<ul><li><u>following:</u></li><li>8. Provider attests the patient has high-risk breast cancer based on which of the following? Check</li></ul>				
all that apply:				
☐ Histopathological tests showing four or more (≥ 4) axillary lymph nodes are affected (pALN N2 or N3 disease)				
☐ Histopathological tests showing one to three axillary lymph nodes are affected ☐ Tumor size is ≥ 5 cm				
Histopathological grade 3 disease (G3)				
☐ Ki-67 score ≥ 20% as determined by an FDA-approved test Other. Specify:				
9. Has the patient undergone surgical resection of the primary tumor? 🗌 Yes 🗌 No				
10. Does the patient have a history of failure or intolerance using one of the following treatment modalities? Check all that apply.				
Radiotherapy				
Taxane (e.g., docetaxel)				
Anthracycline (e.g., doxorubicin) based chemotherapy				
Request for Systemic therapy of recurrent, advanced, or metastatic breast cancer, answer the				
following:				
11.Is the treatment being prescribed as a <u>first-line systemic therapy</u> ?  Yes  No If yes, please select all that apply:				
The patient is a postmenopausal female, premenopausal or perimenopausal female receiving ovarian suppression/ablation (e.g., surgical ablation, suppression with GnRH therapy				
[e.g., leuprolide], etc.) The patient is hormone suppressed male (e.g., GnRH therapy [e.g., leuprolide] used				
concomitantly)				
12. Is the treatment being prescribed as a <u>second-line systemic therapy</u> ?  Yes No				
If yes, please select all that apply: The medication will be used in combination with fulvestrant (Faslodex)				
The patient had disease progression on, or after primary endocrine therapy (as adjuvant or first-line systemic therapy)				
The patient is a postmenopausal female, premenopausal or perimenopausal female receiving ovarian suppression/ablation (e.g., surgical ablation, suppression with GnRH therapy				
[e.g., leuprolide]				
The patient is hormone suppressed male (e.g., GnRH therapy [e.g., leuprolide] used concomitantly)				
13. Is the treatment being prescribed as a <u>subsequent-line (3<sup>rd</sup> line or later) systemic therapy in</u>				
metastatic (stage IV, M1) setting?  Yes No				
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<ul> <li>Patient had disease progression on, or after endocrine therapy AND systemic chemotherapy (not containing a CDK 4/6 inhibitor) in the metastatic setting.</li> <li>The request is for abemaciclib (Verzenio) monotherapy.</li> </ul>				
CHART NOTES & LABS ARE REQUIRED WITH THIS REQUEST				
Prescriber signature	Prescriber specialty	Date		

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