

**US Family Health Plan
Prior Authorization Request Form for
ribociclib (Kisqali) and ribociclib/letrozole (Kisqali Femara Co-Pack)**

To be completed and signed by the prescriber. To be used only for prescriptions which are to be filled through the Department of Defense (DoD) US Family Health Plan Pharmacy Program. US Family Health Plan is a TRICARE contractor for DoD.

The completed form may be faxed to 855-273-5735

OR

The patient may attach the completed form to the prescription and **mail** it to:
Attn: Pharmacy, 77 Warren St, Brighton, MA 02135

QUESTIONS? Call 1-877-880-7007

NOTE: Medical documentation may be requested or your request could be denied.

Step 1 Please complete patient and physician information (please print):

Patient Name: _____ Address: _____ Sponsor ID #: _____ Date of Birth: _____	Physician Name: _____ Address: _____ Phone #: _____ Secure Fax #: _____
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Step 2 Please complete the clinical assessment:

1. Is the requested medication prescribed by or in consultation with an oncologist?	<input type="checkbox"/> Yes Proceed to question 2	<input type="checkbox"/> No STOP Coverage not approved
2. Is the patient currently taking another cyclin-dependent kinase inhibitor?	<input type="checkbox"/> Yes STOP Coverage not approved	<input type="checkbox"/> No Proceed to question 3
3. Is the provider aware and has informed the patient of the risks of neutropenia and interstitial lung disease?	<input type="checkbox"/> Yes Proceed to question 4	<input type="checkbox"/> No STOP Coverage not approved
4. Is the provider aware and has informed the patient of the risk of QT prolongation and hepatobiliary toxicity?	<input type="checkbox"/> Yes Proceed to question 5	<input type="checkbox"/> No STOP Coverage not approved
5. Does the patient have advanced or metastatic hormone receptor (HR(+))/HER2(-) breast cancer?	<input type="checkbox"/> Yes Proceed to question 8	<input type="checkbox"/> No Proceed to question 6
6. Please provide the diagnosis.	_____ Proceed to question 7	
7. Is the diagnosis cited in the National Comprehensive Cancer Network (NCCN) guidelines as a category 1, 2A, or 2B recommendation?	<input type="checkbox"/> Yes Proceed to question 11	<input type="checkbox"/> No STOP Coverage not approved

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8. What is the patient's gender?	<input type="checkbox"/> Female Proceed to question 9	<input type="checkbox"/> Male Proceed to question 21
9. Will the requested medication be used as first-line endocrine therapy in combination with anastrozole, exemestane, or letrozole?	<input type="checkbox"/> Yes Proceed to question 12	<input type="checkbox"/> No Proceed to question 10
10. Will the requested medication be used as first- line or later-line endocrine therapy in combination with fulvestrant?	<input type="checkbox"/> Yes Proceed to question 12	<input type="checkbox"/> No STOP Coverage not approved
11.What is the patient's gender?	<input type="checkbox"/> Female Proceed to question 12	<input type="checkbox"/> Male Proceed to question 21
12. Is the patient of childbearing potential?	<input type="checkbox"/> Yes Proceed to question 13	<input type="checkbox"/> No Proceed to question 17
13. Does the patient agree to use effective contraception during treatment and for at least 3 weeks after cessation of therapy?	<input type="checkbox"/> Yes Proceed to question 14	<input type="checkbox"/> No STOP Coverage not approved
14. Is the patient pregnant?	<input type="checkbox"/> Yes STOP Coverage not approved	<input type="checkbox"/> No Proceed to question 15
15. Has it been confirmed that the patient is not pregnant by negative hCG (human chorionic gonadotropin)?	<input type="checkbox"/> Yes Proceed to question 16	<input type="checkbox"/> No STOP Coverage not approved
16. Will the patient avoid breastfeeding during treatment and for at least 3 weeks after the cessation of treatment?	<input type="checkbox"/> Yes Proceed to question 19	<input type="checkbox"/> No STOP Coverage not approved
17. Is the patient a premenopausal or perimenopausal woman?	<input type="checkbox"/> Yes Proceed to question 18	<input type="checkbox"/> No Proceed to question 19
18. Is the patient receiving ovarian suppression/ablation with a luteinizing hormone-releasing hormone (LHRH) agonist (for example, Lupron [leuprolide], Trelstar [triptorelin], Zoladex [goserelin]), surgical bilateral oophorectomy, or ovarian irradiation?	<input type="checkbox"/> Yes Proceed to question 19	<input type="checkbox"/> No STOP Coverage not approved
19. What is the requested medication?	<input type="checkbox"/> Kisqali Sign and date below	<input type="checkbox"/> Kisqali Femara Proceed to question 20
20. Has the patient been informed of the risk of infertility from letrozole?	<input type="checkbox"/> Yes Sign and date below	<input type="checkbox"/> No STOP Coverage not approved

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21. Has the patient been informed of the risk of infertility?	<input type="checkbox"/> Yes Proceed to question 22	<input type="checkbox"/> No STOP Coverage not approved
22. Is the patient of childbearing potential?	<input type="checkbox"/> Yes Proceed to question 23	<input type="checkbox"/> No Sign and date below
23. Does the patient agree to use effective contraception during treatment and for at least 3 weeks after cessation of therapy?	<input type="checkbox"/> Yes Sign and date below	<input type="checkbox"/> No STOP Coverage not approved

Step
3

I certify the above is true to the best of my knowledge. Please sign and date:

Prescriber Signature

Date

[16 March 2022]