UnitedHealthcare Pharmacy
Clinical Pharmacy Programs

Program Number	2023 P 1260-6
Program	Prior Authorization/Notification
Medication	Kisqali® Femara® Co-Pack (ribociclib/letrozole)
P&T Approval Date	9/2018, 9/2019, 9/2020, 9/2021, 2/2022, 2/2023
Effective Date	5/1/2023; Oxford only: 5/1/2023

1. Background:

Kisqali® Femara® Co-Pack is a co-packaged product containing ribociclib, a kinase inhibitor, and letrozole, an aromatase inhibitor, and is indicated as initial endocrine-based therapy for the treatment of adult patients with hormone receptor (HR)-positive, human epidermal growth factor receptor 2 (HER2)-negative advanced or metastatic breast cancer.

The National Comprehensive Cancer Network (NCCN) recommends the use of Kisqali similarly for men and premenopausal women receiving ovarian ablation/suppression with recurrent unresectable (local or regional) or metastatic HR-positive HER2-negative breast cancer disease in combination with an aromatase inhibitor or fulvestrant. The use of an aromatase inhibitor in men with breast cancer is ineffective without concomitant suppression of testicular steroidogenesis.

Coverage Information:

Members will be required to meet the criteria below for coverage. For members under the age of 19 years, the prescription will automatically process without a coverage review.

Some states mandate benefit coverage for off-label use of medications for some diagnoses or under some circumstances. Some states also mandate usage of other Compendium references. Where such mandates apply, they supersede language in the benefit document or in the notification criteria.

2. Coverage Criteria[a]:

A. <u>Patients less than 19 years of age</u>

1. **Kisqali Femara Co-Pack** will be approved based on the following criterion:

   a. Patient is less than 19 years of age

   Authorization will be issued for 12 months.

B. <u>Breast Cancer</u>

1. **<u>Initial Authorization</u>**

   a. **Kisqali Femara Co-Pack** will be approved based on **<u>all</u>** of the following criteria:

      (1) Diagnosis of advanced, recurrent, or metastatic breast cancer

      -AND-

      (2) Disease is hormone receptor (HR)-positive

-AND-

(3) Disease is human epidermal growth factor receptor 2 (HER2)-negative

Authorization will be issued for 12 months.

2. **Reauthorization**

a. **Kisqali Femara Co-Pack** will be approved based on the following criterion:

(1) Patient does not show evidence of progressive disease while on Kisqali Femara Co-Pack therapy

Authorization will be issued for 12 months.

C. NCCN Recommended Regimens

The drug has been recognized for treatment of the cancer indication by The National Comprehensive Cancer Network (NCCN) Drugs and Biologics Compendium with a Category of Evidence and Consensus of 1, 2A, or 2B.

Authorization will be issued for 12 months.

[a] State mandates may apply. Any federal regulatory requirements and the member specific benefit plan coverage may also impact coverage criteria. Other policies and utilization management programs may apply.

3. Additional Clinical Rules:

- Notwithstanding Coverage Criteria, UnitedHealthcare may approve initial and re-authorization based solely on previous claim/medication history, diagnosis codes (ICD-10) and/or claim logic. Use of automated approval and re-approval processes varies by program and/or therapeutic class.
- Step Therapy and/or Supply limits may be in place.

4. References:

1. Kisqali® Femara® Co-Pack [package insert]. East Hanover, NJ: Novartis Pharmaceuticals Corp. October 2022.
2. The NCCN Drugs and Biologics Compendium (NCCN Compendium™). Available at https://www.nccn.org/professionals/drug_compendium/content/ Accessed January 4, 2023

Program	Prior Authorization/Notification – Kisqali Femara Co-Pack (ribociclib/letrozole)
Change Control	
9/2018	New program.
9/2019	Annual review. Updated background and formatting to align with other CDK 4/6 inhibitor programs. Updated references. Added general NCCN recommended review criteria.
9/2020	Annual review. Minor change to background. Updated references.
9/2021	Annual review with no changes to coverage criteria. Updated

	references.
2/2022	Updated background and references with no change to clinical criteria.
2/2023	Updated background to align with NCCN recommended use. Added state mandate and updated references with no change to clinical criteria.