PREFERRED SPECIALTY MANAGEMENT POLICY

POLICY: Oncology – Cyclin Dependent Kinases 4, 6 Inhibitors Preferred Specialty Management Policy
- Ibrance® (palbociclib capsules and tablets – Pfizer)
- Kisqali® (ribociclib tablets – Novartis)
- Kisqali® Femara® Co-Pack (ribociclib tablets; letrozole tablets, co-packaged – Novartis)
- Verzenio® (abemaciclib tablets – Eli Lilly)

REVIEW DATE: 02/22/2023; selected revision 05/10/2023 and 05/17/2023

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CIGNA NATIONAL FORMULARY COVERAGE:

OVERVIEW

Ibrance, Kisqali/Kisqali Femara Co-Pack, and Verzenio are cyclin-dependent kinase (CDK) 4, 6 inhibitors indicated for use in adults with **hormone receptor positive (HR+), human epidermal growth factor receptor 2 (HER2)-negative advanced or metastatic breast cancer** in the following settings:[1-4]
- All three agents are indicated in combination with an aromatase inhibitor (AI) as initial endocrine-based therapy.
- Ibrance and Verzenio are indicated in combination with fulvestrant for disease progression following endocrine therapy. Kisqali in combination with fulvestrant is approved for use in postmenopausal women or men as initial endocrine-based therapy or following disease progression on endocrine therapy.
- Verzenio is the only agent indicated for use as monotherapy for disease progression following endocrine therapy and prior chemotherapy in the metastatic setting.

- Verzenio is the only agent indicated for use in combination with endocrine therapy (tamoxifen or an AI) for the adjuvant treatment of node-positive, early breast cancer at high risk of recurrence.

Table 1. FDA-Approved Indications for CDK 4, 6 Inhibitors in HR+, HER2-Negative Breast Cancer.[1-4]

	Ibrance® (palbociclib capsules, tablets)	**Kisqali® (ribociclib tablets)**	**Kisqali® Femara® Co-Pack (ribociclib tablets; letrozole tablets, co-packaged)**	**Verzenio® (abemaciclib tablets)**
Early Breast Cancer				
Use in combination with endocrine therapy (AI or tamoxifen) for node positive disease at high risk of recurrence[a]	Not indicated	Not indicated	Not indicated	√
Advanced or Metastatic Breast Cancer				
Use in combination with an AI				
Initial therapy in postmenopausal women[b]	√	√	√	√
Initial therapy in pre/perimenopausal women[b]	√	√	√	√
Initial therapy in men[b]	√	√	√	√

Table 1 (continued). FDA-Approved Indications for CDK 4, 6 Inhibitors in HR+, HER2-Negative Breast Cancer. [1-4]

	Ibrance® (palbociclib capsules, tablets)	**Kisqali® (ribociclib tablets)**	**Kisqali® Femara® Co-Pack (ribociclib tablets; letrozole tablets, co-packaged)**	**Verzenio® (abemaciclib tablets)**
Use in combination with fulvestrant				
Initial therapy in postmenopausal women[c]	Not indicated	√	Not indicated	Not indicated
Initial therapy in men[c]	Not indicated	√	Not indicated	Not indicated
Subsequent therapy in postmenopausal women[c]	√	√	Not indicated	√
Subsequent therapy in premenopausal women[c]	√	Not indicated	Not indicated	√
Subsequent therapy in men[c]	√	√	Not indicated	√
Use as monotherapy[d]	Not indicated	Not indicated	Not indicated	√

CDK 4, 6 – Cyclin-dependent kinase 4 and 6; HR+ – Hormone receptor positive; HER2 – Human epidermal growth factor receptor 2; AI – Aromatase inhibitor; [a] For the adjuvant treatment of adults who have node-positive, early breast cancer at high risk of recurrence; √ – FDA-approved indication; [b] As initial endocrine-based therapy for the treatment of HR+, HER2-negative advanced or metastatic breast cancer; [c] For the treatment of HR+, HER2-negative advanced or metastatic breast cancer with disease progression following endocrine therapy; [d] For the treatment of adult patients with HR+, HER2-negative advanced or metastatic breast cancer with disease progression following endocrine therapy and prior chemotherapy in the metastatic setting.

Guidelines

The National Comprehensive Cancer Network (NCCN) guidelines on breast cancer (version 4.2023 – March 23, 2023) make the following recommendations for recurrent unresectable (local or regional) or Stage IV HR+ and HER2-negative disease in postmenopausal or premenopausal women receiving ovarian ablation or suppression as "Preferred Regimens" for first-line therapy: Kisqali + AI or fulvestrant (category 1); Verzenio + fulvestrant (category 1); Verzenio + AI (category 2A); Ibrance + AI or fulvestrant (category 2A).[5,6] The guidelines state in a footnote that there is controversy on the choice of CDK 4, 6 inhibitors as there are no direct comparative studies between the agents and there are some differences in the study populations in the Phase III randomized studies. The guidelines also state that in Phase III randomized controlled trials, Kisqali + endocrine therapy has shown overall survival benefit in the first-line setting. CDK 4, 6 inhibitor + fulvestrant is Non-Preferred Product as a "Preferred Regimen" for second- and subsequent-line therapy, if CDK 4, 6 inhibitor was not previously used (category 1). The guidelines state that if there is disease progression while on Ibrance, there are limited Phase II data to support the use of Kisqali in the second-line setting.[5] The guidelines state that in Phase III randomized controlled trials, fulvestrant in combination with a CDK 4, 6 inhibitor has shown overall survival benefit in the second-line setting. In this setting, single-agent Verzenio is Non-Preferred Product as a "Useful in Certain Circumstances" (for subsequent treatment) if there is progression on prior endocrine therapy and prior chemotherapy in the metastatic setting (category 2A). For men with breast cancer, the compendium recommends they be treated similarly to postmenopausal women, except that the use of an AI is ineffective without concomitant suppression of testicular steroidogenesis.[6] The guidelines also recommend Verzenio for 2 years as adjuvant therapy in combination with endocrine therapy in patients with HR+, HER2-negative, high risk (i.e., ≥ 4 positive lymph nodes, or 1 to 3 positive lymph nodes with one or more of the following: Grade 3 disease or tumor size ≥ 5 cm) disease (category 2A).

The PALOMA-2 study failed to show an overall survival benefit when Ibrance was combined with letrozole compared with placebo + letrozole in the first-line setting for postmenopausal patients with HR+, HER2-negative advanced breast cancer. Based on an intention-to-treat analysis, the median overall survival was 53.9 months in the Ibrance plus letrozole arm and 51.2 months in the placebo plus letrozole arm; the difference between the arms was not statistically significant. PALOMA-2 met its primary endpoint of improving progression-free survival, but not the secondary endpoint of overall survival.[7]

The MONALEESA-2 study demonstrated a significant overall survival benefit when Kisqali was combined with letrozole in first-line setting compared with placebo + letrozole (median, 63.9 vs. 51.4 months) in postmenopausal patients with HR+, HER2-negative advanced breast cancer.[8] The MONALEESA-7 study also demonstrated a significant overall survival benefit when Kisqali was combined with endocrine therapy in first-line setting compared with placebo + endocrine therapy (median, 58.7 vs. 48.0 months) in pre/perimenopausal patients with HR+, HER2-negative advanced breast cancer.[9]

POLICY STATEMENT

This Preferred Specialty Management program has been developed to encourage the use of Preferred Products. For all medications (Preferred and Non-Preferred), the patient is required to meet the respective standard *Prior Authorization Policy* criteria. The program also directs the patient to try one of the Preferred Products prior to the approval of a Non-Preferred Product. Requests for Non-Preferred Product will also be reviewed using the exception criteria (below). If the patient meets the standard Prior Authorization Policy criteria for Ibrance but has not tried a Preferred Product, a review will be offered for the Preferred Products using the respective standard *Prior Authorization Policy* criteria. All approvals are provided for the duration noted below.

Preferred: Kisqali, Kisqali Femara Co-Pack, Verzenio
Non-Preferred: Ibrance

Oncology – Cyclin Dependent Kinases 4, 6 Inhibitors non-preferred product(s) is(are) covered as medically necessary when the following non-preferred product exception criteria is(are) met. Any other exception is considered not medically necessary.

NON-PREFERRED PRODUCT EXCEPTION CRITERIA

Non-Preferred Product	Exception Criteria
Ibrance	**1. Breast Cancer.** Approve for 1 year if the patient meets the following criteria (A or B): **A)** Patient meets both of the following criteria (i and ii): **i.** Patient meets the standard *Oncology – Ibrance Prior Authorization (PA) Policy* criteria; AND **ii.** Patient meets ONE of the following criteria (a or b): **a)** Patient has been taking Ibrance and is continuing therapy; OR **b)** Patient has tried one of Kisqali, Kisqali Femara Co-Pack, or Verzenio; OR **B)** If the patient has met the standard *Oncology – Ibrance PA Policy* criteria, but has not met the exception criteria above (**Aii**), offer to review for one of the Preferred Products using either the standard *Oncology – Kisqali and Kisqali Femara Co-Pack PA Policy* criteria **or** the *Oncology – Verzenio PA Policy* criteria. **2. Other Conditions**. Approve for 1 year if the patient meets the standard *Oncology – Ibrance PA Policy* criteria.

REFERENCES

1. Ibrance® capsules and tablets [prescribing information]. New York, NY: Pfizer; December 2022.

2. Kisqali® tablets [prescribing information]. East Hanover, NJ: Novartis; October 2022.
3. Kisqali® Femara® Co-Pack tablets [prescribing information]. East Hanover, NJ: Novartis; October 2022.
4. Verzenio® tablets [prescribing information]. Indianapolis, IN: Eli Lilly; March 2023.
5. The NCCN Breast Cancer Clinical Practice Guidelines in Oncology (version 4.2023 – March 23, 2023). © 2023 National Comprehensive Cancer Network. Available at: http://www.nccn.org. Accessed on May 5, 2023.
6. The NCCN Drugs and Biologics Compendium. © 2023 National Comprehensive Cancer Network. Available at: http://www.nccn.org. Search term: ribociclib. Accessed on February 10, 2023.
7. Finn RS, Rugo HS, Dieras V, et al. Overall survival (OS) with first-line palbociclib plus letrozole (PAL+LET) versus placebo plus letrozole (PBO+LET) in women with estrogen receptor–positive/human epidermal growth factor receptor 2–negative advanced breast cancer (ER+/HER2–ABC): analyses from PALOMA-2 [abstract LBA1003]. Presented at: American Society of Clinical Oncology (ASCO) 2022 Annual Meeting; Chicago, IL; June 3–7, 2022.
8. Hortobagyi GN, Stemmer SM, Burris HA, et al. Overall survival with ribociclib plus letrozole in advanced breast cancer. *N Eng J Med*. 2022;386:942-950.
9. Lu Y, Im S, Colleoni M, et al. Updated overall survival of ribociclib plus endocrine therapy versus endocrine therapy alone in pre-and perimenopausal patients with HR+/HER2- advanced breast cancer in MONALEESA-7: a phase III randomized clinical trial. *Clin Cancer Res*. 2022;28(5):851-859.

HISTORY

Type of Revision	Summary of Changes	Review Date
Annual Revision	No changes to the criteria.	01/26/2022
Selected Revision	The requirement that the patient is pre/perimenopausal was removed from the exception criteria, "Patient will be using Kisqali in combination with an aromatase inhibitor as initial endocrine therapy." The requirement of combination use with an aromatase inhibitor for Kisqali Femara CoPack for initial endocrine therapy was removed.	08/10/2022
Update	**12/29/2022**: The overview section and table 1 were updated based on the following removal of wording "postmenopausal women and men" from Ibrance's FDA labeled indication for: hormone receptor positive (HR+), human epidermal growth factor receptor 2 (HER2)-negative advanced or metastatic breast cancer in combination with an aromatase inhibitor as initial endocrine-based therapy.	--
Annual Revision	No criteria changes.	02/22/2023
Update	**03/03/2023:** The overview section was updated due to change in Verzenio's labeling. The following was removed from the indication of early breast cancer "Ki-67 score ≥ 20%, as determined by an FDA approved test." The following was removed from the indication of advanced and metastatic breast cancer in combination with an aromatase inhibitor as initial endocrine-based therapy, "treatment of postmenopausal women and men."	--
Selected Revision	The preferred and non-preferred agents were updated. Ibrance was moved from a preferred agent to a non-preferred agent. Kisqali and Kisqali Femara Co-Pack were moved from non-preferred agent to preferred agent. Verzenio remains a preferred agent. Exception criteria for Kisqali and Kisqali Femara Co-Pack was removed. Exception criteria for Ibrance was added which includes patients who are currently taking Ibrance and patients who have tried one of Kisqali, Kisqali Femara Co-Pack, and Verzenio. Criteria was added that if a patient has met the Oncology –Ibrance Prior	05/10/2023

	Authorization (PA) Policy criteria, but has not met any one of the exception criteria, offer to review for one of the Preferred Products using either the standard Oncology – Kisqali and Kisqali Femara PA Co-Pack Policy criteria or the Oncology – Verzenio PA Policy criteria.	
Selected Revision	**Breast Cancer**: It was clarified that the exception criteria for a patient who has been taking Ibrance and is continuing therapy; or a patient who has tried one of Kisqali, Kisqali Femara Co-Pack, or Verzenio applies to a patient with Breast Cancer. **Other Conditions:** Indications other than Breast Cancer approve for 1 year if the patient meets the standard *Oncology – Ibrance PA Policy* criteria.	05/17/2023