Kadcyla (ado-trastuzumab emtansine)
PAM – 007

Iowa Medicaid Program	Prior Authorization	**Effective Date**	08/15/2013
Revision Number	11	**Last Reviewed**	10/16/2026
Reviewed By	Medicaid Medical Director	**Next Review**	10/15/2027
Approved By	Medicaid Clinical Advisory Committee	**Approved Date**	08/23/2019

Overview

Medication: [1]	ado-trastuzumab emtansine
Brand Name:	Kadcyla®
Pharmacologic Category:	Antineoplastics; human epidermal growth factor receptor 2 (HER2)-targeted antibody and microtubule inhibitor conjugate
FDA-Approved Indication(s):	1. Breast cancer, metastatic (MBC)* – as a single agent, indicated for the treatment of patients with human epidermal growth factor receptor 2 (HER2)-positive, metastatic breast cancer who previously received trastuzumab and a taxane, separately or in combination. Patients should have either: a. received prior therapy for metastatic disease; or b. developed disease recurrence during or within 6 months of completing adjuvant therapy. 2. Early breast cancer (EBC)* – as a single agent, indicated for the treatment of patients with HER2-positive EBC cancer who have residual invasive disease following neoadjuvant taxane and trastuzumab-based treatment. * Select patients for therapy based on FDA-approved companion diagnostic test
How Supplied:	Single-dose vial: 100 mg or 160 mg
Dosage and Administration:	Intravenous (IV) infusion: 3.6 mg/kg every 3 weeks (21-day cycle) until disease progression or unacceptable toxicity (or for up to a total of 14 cycles for patients with early breast cancer)
Benefit Category:	Medical

BOXED WARNING: HEPATOTOXICITY, CARDIAC TOXICITY, EMBRYO-FETAL TOXICITY

- **Hepatotoxicity:** Serious hepatotoxicity has been reported, including liver failure and death in patients treated with Kadcyla®. Monitor serum transaminases and bilirubin prior to initiation of Kadcyla® treatment and prior to each dose. Reduce dose or discontinue Kadcyla® as appropriate in cases of increased serum transaminases or total bilirubin.
- **Cardiac Toxicity:** Kadcyla® administration may lead to reductions in left ventricular ejection fraction (LVEF). Evaluate left ventricular function in all patients prior to and during treatment with Kadcyla®. Withhold treatment for clinically significant decrease in left ventricular function.
- **Embryo-Fetal Toxicity:** Exposure to Kadcyla® during pregnancy can result in embryo-fetal harm. Advise patients of these risks and the need for effective contraception.

Descriptive Narrative

Globally, breast cancer is the most frequently diagnosed malignancy, accounting for over two million cases each year. It is also the leading cause of cancer death in women worldwide. In the United States, breast cancer is the most common female cancer, and the second most common cause of cancer death in women.[2] An estimated 3,030 new cases of breast cancer in females will be diagnosed in Iowa in 2026, making it the second most commonly diagnosed cancer in the state (second only to prostate cancer), and with an estimated 370 deaths, it is the fourth highest cause of cancer deaths in Iowa in 2026.[3]

Up to 5 percent of women diagnosed with breast cancer in the U.S. have metastatic disease at the time of first presentation, despite the gains in early detection, and up to 30 percent of women with early-stage non-metastatic breast cancer at diagnosis will develop distant metastatic disease.

Breast cancer is a heterogenous, phenotypically diverse disease composed of several biologic subtypes that have distinct behavior. Amplification or overexpression of the human epidermal growth factor receptor 2 *(HER2)* oncogene* is present in approximately 15 percent of primary invasive breast cancers. Women with both early-stage and metastatic breast cancer** that meet criteria for HER2 positivity are treated with regimens including HER2-directed therapy.[4]

* *HER2* is a growth-promoting protein on the outside of all breast cells. Breast cancer cells with higher-than-normal levels of HER2 are referred to as HER2-positive.

** Metastatic breast cancer (MBC). Also called Stage IV, MBC is breast cancer that has spread to another part of the body, commonly the liver, brain, bones, or lungs.

Guidelines

The National Comprehensive Cancer Network (NCCN) Clinical Practice Guidelines in Oncology document evidence-based, consensus-driven management to ensure that all patients receive preventive, diagnostic, treatment, and supportive services that are most likely to lead to optimal outcomes. The guidelines are developed and updated by 63 individual panels, comprising over 1,900 clinicians and oncology researchers from the 33 NCCN Member Institutions. The categories for recommendations are based on both the level of clinical evidence available and the degree of consensus within the NCCN Guidelines Panel.

The library of NCCN Guidelines® currently apply to more than 97 percent of people living with cancer or anyone at risk for a diagnosis of cancer in the United States. The guidelines incorporate real-time updates in keeping with the rapid advancements in the field of cancer research and management and are intended to assist all individuals who impact decision-making in cancer care, including physicians, nurses, pharmacists, payers, patients and their families, and others.

The NCCN Guidelines® provide recommendations based on the best evidence available at the time they are derived. Because new data are published continuously, it is essential that the NCCN Guidelines also be continuously updated and revised* to reflect new data and clinical information that may add to or alter current clinical practice standards.[5,6]

The information referenced at the time of this policy writing/revision (note version number and effective date) is from the NCCN Guidelines® for:[7]

- Breast Cancer (v.6.2026 – July 29, 2026)

* To view the most recent and complete version of the guidelines, go online to NCCN.org. NCCN makes no warranties of any kind whatsoever regarding their content, use or application and disclaims any responsibility for their application or use in any way.

Criteria

Prior authorization is required.

HER2-Positive Early Breast Cancer with Residual Disease – Initial Criteria

Kadcyla® is considered medically necessary when **ALL** of the following are met:

1. Diagnosis of human epidermal growth factor receptor 2 *(HER2)*-positive early breast cancer (EBC); **AND**
2. Member has residual invasive disease following neoadjuvant taxane (paclitaxel or docetaxel) and trastuzumab-based treatment; **AND**
3. Kadcyla® will be administered as a single agent treatment regimen; **AND**
4. Member is 18 years of age or older; **AND**
5. Prescribed by, or in consultation with, an oncologist; **AND**
6. Request meets one of the following (a or b):
    a. Regimen prescribed does not exceed 3.6 mg/kg every 3 weeks (21-day cycle) for a maximum of 14 cycles of therapy; or,
    b. Regimen is supported by clinical practice guidelines (i.e., must be recommended in NCCN Guidelines®). Supporting clinical documentation must be provided with any request for which regimen prescribed does not align with FDA-approved labeling.

HER2-Positive Metastatic Breast Cancer – Initial Criteria

Kadcyla® is considered medically necessary when **ALL** of the following are met:

1. Diagnosis of human epidermal growth factor receptor 2 *(HER2)*-positive metastatic breast cancer (MBC); **AND**
2. Member previously treated with trastuzumab and a taxane (paclitaxel or docetaxel), separately or in combination; **AND**
3. Member meets **ONE** of the following (a or b):
    a. Received prior therapy for metastatic disease; **OR**
    b. Developed disease recurrence during or within 6 months of completing adjuvant therapy; **AND**
4. Kadcyla® will be administered as a single agent treatment regimen; **AND**
5. Member is 18 years of age or older; **AND**
6. Prescribed by, or in consultation with, an oncologist; **AND**
7. Request meets one of the following (a or b):
    a. Regimen prescribed does not exceed 3.6 mg/kg every 3 weeks (21-day cycle); or,
    b. Regimen is supported by clinical practice guidelines (i.e., must be recommended in NCCN Guidelines®). Supporting clinical documentation must be provided with any request for which regimen prescribed does not align with FDA-approved labeling.

Continuation Criteria (all above indications)

Kadcyla® is considered medically necessary for continuation of therapy when **ALL** of the following are met:

1. Member is currently receiving medication through the Iowa Medicaid benefit or has previously met initial approval criteria; **AND**
2. Documentation of positive clinical response to therapy, as demonstrated by tumor response or lack of disease progression, and an acceptable toxicity profile; **AND**
3. For early breast cancer (EBC) **ONLY**: Member has not received the maximum of 14 cycles of treatment with Kadcyla®; **AND**
4. Prescribed by, or in consultation with, an oncologist; **AND**
5. Request meets one of the following (a or b):
    a. Regimen prescribed does not exceed 3.6 mg/kg every 3 weeks (21-day cycle):
        1) for a maximum of 14 cycles of therapy for early breast cancer; or
        2) until disease progression or unacceptable toxicity in metastatic breast cancer; or
    b. Regimen is supported by clinical practice guidelines (i.e., must be recommended in NCCN Guidelines®). Supporting clinical documentation must be provided with any request for which regimen prescribed does not align with FDA-approved labeling.

Approval Duration and Quantity Limits

Indication	**Early Breast Cancer (EBC)**	**Metastatic Breast Cancer (MBC)**
Duration of Authorization	**Initial:** 6 months or 8 cycles, whichever comes first	**Initial:** 6 months or 8 cycles, whichever comes first
	Continuation: 4 months (6 cycles), or until maximum 14 cycles are completed	**Continuation:** 12 months or 17 cycles whichever comes first
Quantity Limits	• 3.6 mg/kg every 3 weeks (21-day cycle) • Treat for a total of 14 cycles unless there is disease recurrence or unmanageable toxicity	• 3.6 mg/kg every 3 weeks (21-day cycle) • Treat until disease progression or unmanageable toxicity

Coding and Product Information

The following list(s) of codes and product information are provided for reference purposes only and may not be all inclusive. The inclusion of a code does not imply any right to reimbursement or guarantee claim payment, nor does the exclusion of a code imply that its association to the HCPCS code is inappropriate.

HCPCS	**Description**
J9354	Injection, ado-trastuzumab emtansine, 1 mg

ICD-10	**Description**
C50.11-C50.929	Malignant neoplasm of breast [HER2 positive] [not covered if HER2 negative]
C79.81	Secondary malignant neoplasm of breast
D05.00-D05.92	Carcinoma in situ of breast

NDC (Strength)	**Labeler**	**Dosage**	**Pkg Size**	**Pkg Qty**	**Units/ Pkg**
50242-0087-01 (160 mg)	Genentech, Inc. (50242)	1 mg	1	EA	160
50242-0088-01 (100 mg)	Genentech, Inc. (50242)	1 mg	1	EA	100

Compliance

1. Should conflict exist between the policy and applicable statute, the applicable statute shall supersede.
2. Federal and State law, as well as contract language, including definitions and specific contract provisions or exclusions, take precedence over medical policy and must be considered first in determining eligibility for coverage.
3. Medical technology is constantly evolving, and Iowa Medicaid reserves the right to review and update medical policy on an annual or as-needed basis.

Medical necessity guidelines have been developed for determining coverage for member benefits and are published to provide a better understanding of the basis upon which coverage decisions are made. Medical necessity guidelines are developed for selected physician-administered medications found to be safe and proven to be effective in a limited, defined population or clinical

circumstances. They include concise clinical coverage criteria based on current literature review, consultation with practicing physicians in the service area who are medical experts in the particular field, FDA and other government agency policies, and standards adopted by national accreditation organizations. Criteria are revised and updated annually, or more frequently if new evidence becomes available that suggests needed revisions.

References

[1] Kadcyla® prescribing information (03/2026). Genentech, Inc.: San Francisco, CA. Available online: www.kadcyla-hcp.com. Accessed July 29, 2026.

[2] Joe BN. Clinical features, diagnosis, and staging of newly diagnosed breast cancer. Vora SR, ed. UpToDate. Waltham, MA: UpToDate Inc. www.uptodate.com. Accessed July 29, 2026.

[3] American Cancer Society: Cancer Statistics Center. State of Iowa. Available online at cancerstatisticscenter.cancer.org/states/iowa. Accessed October 16, 2026.

[4] Yamauchi H, Bleiweiss IJ. HER2 and predicting response to therapy in breast cancer. Vora SR, ed. UpToDate. Waltham, MA: UpToDate Inc. www.uptodate.com. Accessed July 29, 2026.

[5] National Comprehensive Cancer Network (NCCN). Guidelines Process: About Clinical Practice Guidelines. Available online at www.nccn.org. Accessed October 20, 2025.

[6] National Comprehensive Cancer Network (NCCN). Guidelines Process: Development and Update of Guidelines. Available online at www.nccn.org. Accessed October 20, 2025.

[7] NCCN Clinical Practice Guidelines in Oncology. The NCCN Guidelines® are continuously updated and revised to reflect new data and clinical information that may add to or alter current clinical practice standards. To view the most recent and complete version, go online to NCCN.org. NCCN Guidelines® referenced at time of this revision (note version number and effective date):

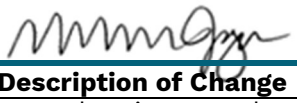
- Breast Cancer (v.6.2026 – July 29, 2026)

Development of utilization management criteria may also involve research into other state Medicaid programs, other payer policies, consultation with experts and review by the Medicaid Clinical Advisory Committee (CAC). These sources may not be referenced individually unless they are specifically published and are otherwise applicable to the criteria at issue.

Criteria Change History

Change Date	Changed By	Description of Change	Version
[mm/dd/yyyy]	CAC		
Signature			

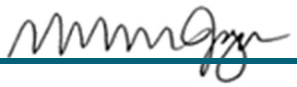
Change Date	Changed By	Description of Change	Version
[mm/dd/yyyy]	CAC		
Signature			

Change Date	Changed By	Description of Change	Version
10/16/2026	CAC	Annual review. Updated ACS statistics for 2026 (website was not available at time of initial review). Updated references. Policy format update: Removed current NCCN Guidelines recommendations, as guidelines may change more frequently than the review cadence of this document. Will continue to reference NCCN when writing criteria, and include the specific version referenced at the time of the policy writing/revision.	11
Signature William (Bill) Jagiello, DO			

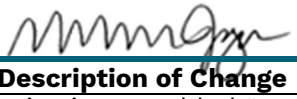
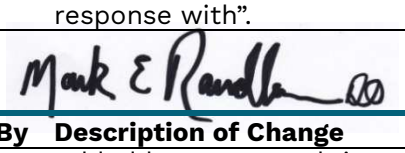
Change Date	Changed By	Description of Change	Version
10/17/2025	CAC	Annual review. No changes.	10
Signature William (Bill) Jagiello, DO			

Change Date	Changed By	Description of Change	Version
10/18/2024	CAC	Annual review. Reviewed NCCN Guidelines, updated where appropriate. Updated references.	9
Signature William (Bill) Jagiello, DO			

Change Date	Changed By	Description of Change	Version
10/20/2023	CAC	Annual review. Added boxed warning to Overview section (hepatotoxicity, cardiac toxicity, embryo-fetal toxicity). Updated NCCN Guidelines. Added dosing regimens into clinical criteria.	8
Signature William (Bill) Jagiello, DO			

Change Date	Changed By	Description of Change	Version
10/21/2022	CAC	Annual review. Updated NCCN recommendations. Added "Member is 18 years of age or older" to initial coverage criteria. Added standard language to continuation criteria: "1. Member is currently receiving medication through the Iowa Medicaid benefit or has previously met initial approval criteria." Updated references. Prior authorization requirement re-implemented effective 6/1/22 to align with current policy.	7
Signature William (Bill) Jagiello, DO			

Criteria Change History *(continued)*

Change Date	Changed By	Description of Change	Version
10/15/2021	CAC	Annual review.	6
Signature			
William (Bill) Jagiello, DO			
Change Date	**Changed By**	**Description of Change**	**Version**
07/17/2020	CAC	Prior authorization requirement removed.	5
Signature			
William (Bill) Jagiello, DO			
Change Date	**Changed By**	**Description of Change**	**Version**
07/16/2015	Medical Director	Criterion #1 added "over expression of the HER2 gene". Criterion #2 added "failed treatment or shown inadequate response with".	4
Signature			
Mark E. Randleman, DO			
Change Date	**Changed By**	**Description of Change**	**Version**
07/17/2015	CAC	Added last paragraph in References.	3
Signature			
Change Date	**Changed By**	**Description of Change**	**Version**
07/14/2015	Medical Director	Updated NCCN reference.	2
Signature			
Change Date	**Changed By**	**Description of Change**	**Version**
07/18/2014	CAC	Removed narrative that was a duplication of what is listed under criteria.	1
Signature			

CAC = Medicaid Clinical Advisory Committee