

Reference number(s)
5608-D

This document applies to the following:

Formulary	Applies
Standard Control (SF)	<input type="checkbox"/>
Standard Control – Choice (SCCF)	<input type="checkbox"/>
Preferred Drug Plan Design (PDPD)	<input type="checkbox"/>
Advanced Control Specialty (ACSF)	<input type="checkbox"/>
Advanced Control Specialty – Choice (ACSCF)	<input type="checkbox"/>
Managed Medicaid Template (MMT)	<input type="checkbox"/>
Marketplace (MF)	<input type="checkbox"/>
Aetna Small Group Affordable Care Act (SG ACA)	<input type="checkbox"/>
Aetna Health Exchange (AHE)	<input type="checkbox"/>
Aetna Individual Lives (IVL)	<input type="checkbox"/>
Value (VF)	<input type="checkbox"/>

Formulary	Applies
New to Market (NTM)	<input type="checkbox"/>
Standard Formulary Chart (SFC)	<input type="checkbox"/>
Basic Control Chart Preferred Drug Plan Design (BCC PDPD)	<input type="checkbox"/>
Advanced Control Specialty Formulary Chart (ACSF)	<input type="checkbox"/>
Value Formulary Chart (VFC)	<input type="checkbox"/>
Medical Benefit	<input checked="" type="checkbox"/>
Medical Benefit: Advanced Biosimilars First	<input type="checkbox"/>
Medical Benefit: Managed Medicaid (MMMB)	<input type="checkbox"/>
Medicare Part B	<input type="checkbox"/>
Medicare Part B: Advanced Biosimilars First	<input type="checkbox"/>

Exceptions Criteria

Breast Cancer

This document informs prescribers of preferred products and provides an exception process for targeted products through prior authorization.

These criteria were developed to align with the following: Medical Benefit.

Plan Design Summary

This program applies to the breast cancer products specified in this document. Coverage for targeted products is provided based on clinical circumstances that would exclude the use of the preferred products and may be based on previous use of a product. The coverage review process will ascertain situations where a clinical exception can be made. This program applies to members who are new to treatment with a targeted product for the first time.

Each referral is reviewed based on all utilization management (UM) programs implemented for the client.

Table. HER2-targeted antibodies

Medications considered formulary or preferred on your plan may still require a clinical prior authorization review.

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	Products
Preferred	<ul style="list-style-type: none"> • Enhertu (fam-trastuzumab deruxtecan-nxki) • Kadcyla (ado-trastuzumab emtansine) • Perjeta (pertuzumab) • Phesgo (pertuzumab, trastuzumab, and hyaluronidase-zzxf)
Target	<ul style="list-style-type: none"> • Margenza (margetuximab-cmkb)

Exception Criteria

This program applies to members requesting treatment for breast cancer.

Coverage for the targeted product is provided when either of the following criteria is met:

- Member is currently receiving treatment with the targeted product, excluding when the requested targeted product is obtained as samples or via manufacturer’s patient assistance programs.
- Member has a documented inadequate response or intolerable adverse event with at least three of the preferred products.

References

1. Enhertu [package insert]. Basking Ridge, NJ: Daiichi Sankyo, Inc.; November 2022.
2. Kadcyla [package insert]. South San Francisco, CA: Genentech, Inc.; February 2022.
3. Margenza [package insert]. Rockville, MD: MacroGenics, Inc.; May 2023.
4. Perjeta [package insert]. South San Francisco, CA: Genentech, Inc.; February 2021.
5. Phesgo [package insert]. South San Francisco, CA: Genentech, Inc.; June 2020.