

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 checked="" type="checkbox"/>

Formulary	Applies
New to Market (NTM)	<input type="checkbox"/>
Standard Formulary Chart (SFC)	<input checked="" type="checkbox"/>
Basic Control Chart Preferred Drug Plan Design (BCC PDPD)	<input type="checkbox"/>
Advanced Control Specialty Formulary Chart (ACSFC)	<input checked="" type="checkbox"/>
Value Formulary Chart (VFC)	<input checked="" type="checkbox"/>
Medical Benefit	<input 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

Osteoporosis

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: Value Formulary (VF), Standard Formulary Chart (SFC), Advanced Control Specialty Formulary Chart (ACSFC), and Value Formulary Chart (VFC).

Plan Design Summary

This program applies to the osteoporosis products specified in this document. Coverage for the 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 initiating a new treatment regimen with Evenity. This program also applies to all members requesting treatment with Forteo.

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

Table. Osteoporosis Products

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

	Product(s)
Preferred	<ul style="list-style-type: none"> • Prolia (denosumab) • teriparatide (generic) • Tymlos (abaloparatide)
Target	<ul style="list-style-type: none"> • Evenity (romosozumab-aqqg) • Forteo (teriparatide)

Exception Criteria

This program applies to members requesting treatment for an indication that is FDA-approved for any of the preferred products.

Evenity

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

- There is documentation that the member is currently undergoing treatment with the targeted product Evenity, and coverage is required to complete the current course of treatment.
- Member has a documented inadequate response, intolerable adverse event, or contraindication to all of the preferred products (e.g., cumulative treatment with Tymlos exceeding 24 months in a patient's lifetime).

Forteo

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

- Member will be using the targeted product Forteo for treatment of glucocorticoid-induced osteoporosis and has had a documented intolerable adverse event to the preferred product generic teriparatide, and the adverse event was not an expected adverse event attributed to the active ingredient as described in the prescribing information.
- Member meets both of the following criteria:
 - Member has had a documented intolerable adverse event to the preferred product generic teriparatide, and the adverse event was not an expected adverse event attributed to the active ingredient as described in the prescribing information.
 - Member has a documented inadequate response, intolerable adverse event, or contraindication to the preferred product Tymlos (e.g., cumulative treatment with Tymlos exceeding 24 months in a patient's lifetime).

Reference number(s)
6627-D

References

1. Evenity [package insert]. Thousand Oaks, CA: Amgen, Inc.; April 2024.
2. Forteo [package insert]. Indianapolis, IN: Lilly USA, LLC; July 2024.
3. Prolia [package insert]. Thousand Oaks, CA: Amgen Inc.; March 2024.
4. Teriparatide [package insert]. Weston, FL: Apotex Corp.; October 2024.
5. Tymlos [package insert]. Boston, MA: Radius Health, Inc.; December 2023.