



## **BONE DENSITY REGULATORS PRIOR AUTHORIZATION FORM** (form effective 1/5/2026)

FAX this completed form to (844) 205-3386

OR Mail requests to: Pharmacy Department | 5 River Park Place East, Suite 210 | Fresno, CA 93720

OR Prior authorization may be completed at <https://www.covermymeds.com/main/prior-authorization-forms/>

Prior authorization guidelines **Bone Density Regulators** and **Quantity Limits/Daily Dose Limits** are available on the PA Health & Wellness website at <https://www.pahealthwellness.com/providers/pharmacy.html>.

<input type="checkbox"/> New request	<input type="checkbox"/> Renewal request	Total pages: _____	Prescriber name:	
Name of office contact:		Specialty:		
Contact's phone number:		NPI:	State license #:	
LTC facility contact/phone:		Street address:		
Member name:		City/state/zip:		
Member ID#:	DOB:	Phone:	Fax:	

### **CLINICAL INFORMATION**

Drug requested:	Strength:	Dosage form:	
Dose/directions:		Quantity:	Refills:
Diagnosis ( <u>submit documentation</u> ):		Dx code ( <u>required</u> ):	

**Complete all sections that apply to the member and this request.**

***Check all that apply and submit documentation for each item.***

#### **INITIAL requests**

##### **1. For treatment of an OSTEOPOROSIS-RELATED condition:**

Has results of a recent bone mineral density test → Document T-score: \_\_\_\_\_ Date of test: \_\_\_\_\_

Was evaluated for other possible causes of osteoporosis and has results of the following lab tests:

<input type="checkbox"/> CBC	<input type="checkbox"/> Phosphorous	<input type="checkbox"/> Total protein	<input type="checkbox"/> Thyroid stimulating hormone (TSH)
<input type="checkbox"/> Vitamin D	<input type="checkbox"/> Creatinine	<input type="checkbox"/> Urinary calcium excretion	<input type="checkbox"/> Intact parathyroid hormone (PTH)
<input type="checkbox"/> Ionized calcium	<input type="checkbox"/> Albumin	<input type="checkbox"/> Testosterone (if male)	<input type="checkbox"/> Liver enzymes (specifically alkaline phosphatase)



**2. For an ANABOLIC AGENT (e.g., Bonsity, Evenity, Forteo, teriparatide):**

- Has a history of fragility fracture
- Has a history of multiple vertebral fractures
- Has a history of trial and failure of or a contraindication or an intolerance to bisphosphonates
- Request will not exceed the cumulative treatment duration recommended in the FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature

**For a PARATHYROID HORMONE ANALOG (e.g., abaloparatide [Tymlos], teriparatide [e.g., Bonsity, Forteo])** – check all that apply to the member:

- Paget's disease of the bone
- Metabolic bone disease other than osteoporosis
- Bone metastases
- Hypercalcemic disorder(s)
- History of skeletal malignancies
- Unexplained elevations of alkaline phosphatase
- Open epiphyses
- Prior external beam or implant radiation therapy involving the skeleton

**For EVENITY** – check all that apply to the member:

- History of myocardial infarction
- History of stroke

**For EVENITY or TYMLOS:**

- Has a contraindication or an intolerance to teriparatide

**For FORTEO and BONSITY:**

- Has a contraindication or an intolerance to generic teriparatide that would not be expected to occur with the requested drug

**3. For EVISTA (raloxifene):**

- Check all that apply to the member:
  - History of venous thromboembolic events (including deep vein thrombosis, pulmonary embolism, and retinal vein thrombosis)
  - History of breast cancer
- Has ONE or more risk factors for stroke:
  - History of stroke or TIA
  - Hypertension
  - other: \_\_\_\_\_
  - Atrial fibrillation
  - Cigarette smoker
- If member has one or more risk factors for stroke, was counseled by the prescriber about the increased risk of death due to stroke
- Is a post-menopausal or post-oophorectomy female
- Is at high risk for fracture defined by at least ONE of the following:
  - A 10-year probability of hip fracture  $\geq 3\%$  based on the US-adapted WHO algorithm
  - A 10-year probability of major fracture related to osteoporosis  $\geq 20\%$  based on the US-adapted WHO algorithm
  - A history of fragility fracture of the proximal humerus, pelvis, or distal forearm
  - A history of low-trauma spine or hip fracture
- Is at high risk for invasive breast cancer defined by at least ONE of the following:
  - Prior biopsy with lobular carcinoma in situ (LCIS) or atypical hyperplasia
  - One or more first-degree relatives with breast cancer
  - A 5-year predicted risk of breast cancer  $\geq 1.66\%$  (based on the modified Gail model)
- Has a history of trial and failure of or a contraindication or an intolerance to oral bisphosphonates

**4. For DENOSUMAB 120 MG/1.7 ML (i.e., Xgeva and corresponding biosimilars), the member is being treated for at least ONE of the following:**

- Bone metastases from solid tumors



- Giant cell tumor of the bone
- Hypercalcemia of malignancy
- Multiple myeloma
- A diagnosis not in the list above that is supported by FDA-approved package labeling, peer-reviewed medical literature, or nationally recognized medical compendia

**5. For ALL OTHER Bone Density Regulators:**

- Is at high risk for fracture defined by at least ONE of the following:
  - A 10-year probability of hip fracture  $\geq 3\%$  based on the US-adapted WHO algorithm
  - A 10-year probability of major fracture related to osteoporosis  $\geq 20\%$  based on the US-adapted WHO algorithm
  - A history of fragility fracture of the proximal humerus, pelvis, or distal forearm
  - A history of low-trauma spine or hip fracture
- Has a history of trial and failure of or a contraindication or an intolerance to the preferred Bone Density Regulators (Refer to <https://papdl.com/preferred-drug-list> for a list of preferred and non-preferred drugs in this class.)
- For a PARENTERAL bisphosphonate:
  - Has a contraindication or an intolerance to oral bisphosphonates

**RENEWAL requests**

**1. For ALL renewal requests:**

- The member's condition has stabilized since starting the requested medication
- The member continues to benefit from the requested medication

**PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION TO 844-205-3386**

Prescriber Signature:	Date:
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Pharmacy Department will respond via fax or phone within 24 hours.

Requests for prior authorization (PA) requests must include member name, ID#, and drug name. Please include lab reports with requests when appropriate (e.g., Culture and Sensitivity; Hemoglobin A1C; Serum Creatinine; CD4; Hematocrit; WBC, etc.)