



Commonwealth of Massachusetts
MassHealth Drug Utilization Review Program
 P.O. Box 2586, Worcester, MA 01613-2586
Fax: (877) 208-7428 **Phone:** (800) 745-7318

Osteoporosis Agents and Calcium Regulators Prior Authorization Request

MassHealth reviews requests for prior authorization (PA) on the basis of medical necessity only. If MassHealth approves the request, payment is still subject to all general conditions of MassHealth, including current member eligibility, other insurance, and program restrictions. MassHealth will notify the requesting provider and member of its decision. Keep a copy of this form for your records. If faxing this form, please use black ink.

Additional information about these agents, including PA requirements and preferred products, can be found within the MassHealth Drug List at www.mass.gov/druglist.

Member information

Last name _____ First name _____ MI _____
 MassHealth member ID # _____ Date of birth _____
 Gender (Check one.) F M Member's place of residence home nursing facility

Medication information

Medication requested

Bisphosphonates

- | | |
|--|--|
| <input type="checkbox"/> Fosamax Plus D (alendronate/cholecalciferol) (Section VI) | <input type="checkbox"/> risedronate (Section I) |
| <input type="checkbox"/> ibandronate IV (Section I) | <input type="checkbox"/> risedronate delayed-release (Section I) |

Miscellaneous Agents

- | | |
|---|---|
| <input type="checkbox"/> calcitonin salmon injection (Section III) | <input type="checkbox"/> Prolia (denosumab) (Section I) |
| <input type="checkbox"/> Evenity (romosozumab-aqqg) (Section III) | <input type="checkbox"/> teriparatide (Section I and V) |
| <input type="checkbox"/> Forteo (teriparatide) (Section I) | <input type="checkbox"/> Tymlos (abaloparatide) (Section III) |
| <input type="checkbox"/> Natpara (parathyroid hormone) (Section IV) | <input type="checkbox"/> Xgeva (denosumab) (Section VII and VIII) |

Dose, frequency, and duration of medication requested _____

Indication (Check all that apply.)

- | | |
|---|---|
| <input type="checkbox"/> Giant cell tumor of the bone (Xgeva) (Section VIII) | <input type="checkbox"/> Prevention of bone loss in men receiving androgen deprivation therapy for prostate cancer |
| <input type="checkbox"/> Glucocorticoid-Induced Osteoporosis (GIO) (Section II) | <input type="checkbox"/> Prevention of bone loss in women receiving aromatase inhibitors for breast cancer |
| <input type="checkbox"/> Hypercalcemia | <input type="checkbox"/> Prevention of skeletal-related events secondary to bone metastases in cancer related to solid tumors (Xgeva) (Section VII) |
| <input type="checkbox"/> Hypercalcemia of malignancy (Xgeva) (Section VII) | <input type="checkbox"/> Prevention of skeletal-related events secondary to multiple myeloma (Xgeva) (Section VII) |
| <input type="checkbox"/> Hypocalcemia with hypoparathyroidism | <input type="checkbox"/> Primary/Hypogonadal Osteoporosis |
| <input type="checkbox"/> Osteopenia | <input type="checkbox"/> Other _____ |
| <input type="checkbox"/> Paget's Disease | |
| <input type="checkbox"/> Postmenopausal Osteoporosis (PMO) | |

Please indicate billing preference. Pharmacy Prescriber in-office Hospital outpatient

For hospital outpatient billing, provide department-specific facility NPI. _____

Drug NDC (if known) or service code _____

Section I. Please complete for bisphosphonates, Forteo, Prolia, and teriparatide as indicated above.

1. Please provide results of bone mineral density (BMD) measurements (T-scores of total hip and lumbar vertebrae). _____
2. Has the member had a radiographically confirmed fracture?
 Yes. Please provide site and date below.
Site _____ Date _____
 No
3. Please list all non-modifiable risk factors for fracture in this member.

4. Has the member tried an oral bisphosphonate and experienced an adverse reaction or inadequate response?
 Yes. Please list the dates/duration of oral bisphosphonate trial and outcomes in Section IX below.*
 No. Please document if there is a contraindication to oral bisphosphonates.

5. If the request is for Forteo or teriparatide, has the member tried Prolia or an intravenous bisphosphonate and experienced an adverse reaction or inadequate response?
 Yes. Please list the drug names, dates/duration of trials and outcomes in Section IX below.*
 No. Please document if there is a contraindication to Prolia and intravenous bisphosphonates.

Section II. Please complete for all agents being requested for the treatment or prevention of Glucocorticoid-Induced Osteoporosis (GIO).

Please provide specifics of the member's chronic glucocorticoid use.

Drug _____ Dose and Frequency _____ Dates/Duration _____

Section III. Please complete for calcitonin salmon injection, Evenity, and Tymlos requests.

Please attach supporting documentation of the diagnosis, BMD measurements, medical necessity for the requested agent, fracture risk factors, and previous trials including oral bisphosphonates, IV bisphosphonates (ibandronate, pamidronate, zoledronic acid 5 mg), or Prolia as applicable. For calcitonin salmon injection, Evenity, and Tymlos requests, please also attach supporting documentation of previous trials including Forteo as applicable.

Section IV. Please complete for Natpara requests.

Please attach supporting documentation of diagnosis and previous trials including calcium and active vitamin D supplementation as applicable. Please also attach consultation notes from an endocrinologist addressing the use of the requested agent.

Section V. Please complete for teriparatide requests.

Please provide medical necessity for the use of teriparatide instead of Forteo. _____

Section VI. Please complete for Fosamax Plus D requests.

Please provide compelling clinical rationale for why the combination alendronate/cholecalciferol would offer a therapeutic advantage over the individual agents. _____

Section VII. Please complete for Xgeva requests for a diagnosis of prevention of skeletal-related events secondary to bone metastases in cancer related to solid tumors, prevention of skeletal-related events secondary to multiple myeloma, and hypercalcemia of malignancy.

Please indicate prescriber specialty below.

Hematology Oncology Orthopedic Specialist Other _____

If prescriber is not a specialist, please attach consult notes from specialist.

Section VIII. Please complete for Xgeva requests for a diagnosis of giant cell tumor of the bone.

Please describe surgical history and/or prognosis. If surgery is not appropriate for this member, please explain.

Section IX. Please complete for all requests as needed.

Please provide the following information regarding previous trials.*

Drug name/Therapy _____ Dates/duration of use _____

Did the member experience any of the following? Adverse reaction Inadequate response

Briefly describe details of adverse reaction or inadequate response.

Drug name/Therapy _____ Dates/duration of use _____

Did the member experience any of the following? Adverse reaction Inadequate response

Briefly describe details of adverse reaction or inadequate response.

* Please attach a letter documenting additional trials as necessary.

Section X. Please complete for all requests for non-preferred drug products if one or more preferred drug products have been designated for this class of drugs.

If one or more preferred drug products have been designated for this class of drugs, and if you are requesting PA for a non-preferred drug product, please provide medical necessity for prescribing the non-preferred drug product rather than the preferred drug product.

Prescriber information

Last name* _____ First name* _____ MI _____

NPI* _____ Individual MH Provider ID _____

DEA No. _____ Office Contact Name _____

Address _____ City _____ State _____ Zip _____

E-mail address _____

Telephone No.* _____ Fax No.* _____

* Required

Prescribing provider's attestation, signature, and date

I certify under the pains and penalties of perjury that I am the prescribing provider identified in the Prescriber information section of this form. Any attached statement on my letterhead has been reviewed and signed by me. I certify that the medical necessity information (per 130 CMR 450.204) on this form is true, accurate, and complete, to the best of my knowledge. I understand that I may be subject to civil penalties or criminal prosecution for any falsification, omission, or concealment of any material fact contained herein.

Prescribing provider's signature (Signature and date stamps, or the signature of anyone other than the provider, are not acceptable.)

Signature required _____

Printed name of prescribing provider _____ Date _____