

## INJECTABLE BONE RESORPTION SUPPRESSION AGENTS PRIOR AUTHORIZATION FORM

- To review the prior authorization guidelines for Injectable Bone Resorption Suppression Agents, please refer to the Medical Assistance Prior Authorization of Pharmaceutical Services Handbook Chapter – **Bone Resorption Suppression Agents** (accessible at: <http://www.dhs.pa.gov/provider/pharmacyservices/drugsrequiringclinicalpriorauthorization/index.htm>).
- These agents are also subject to quantity limits. If the requested quantity exceeds the limit, please submit supporting chart documentation (refer to **Quantity Limits / Daily Dose Limits** at: <http://www.dhs.pa.gov/provider/pharmacyservices/quantitylimitslist/index.htm>).

PRIOR AUTHORIZATION REQUEST INFORMATION		PRESCRIBER INFORMATION	
<input type="checkbox"/> New request	<input type="checkbox"/> Additional info	# of pages in request: _____	
<input type="checkbox"/> Renewal request	(PA# _____)	Prescriber name: _____	
Name of office contact: _____		Specialty: _____	
Contact's phone number: _____		State license #: _____	
LTC facility contact/phone: _____		NPI: _____	MA Provider ID#: _____
RECIPIENT INFORMATION		Street address: _____	
Recipient Name: _____		Suite #: _____	City/State/Zip: _____
Recipient ID#: _____	DOB: _____	Phone: _____	Fax: _____

### CLINICAL INFORMATION

<b>Non-preferred injectable medication requested</b>	<input type="checkbox"/> Boniva injection <input type="checkbox"/> Forteo injection* (*please use "Forteo Form") <input type="checkbox"/> ibandronate injection	<input type="checkbox"/> Miacalcin injection <input type="checkbox"/> Prolia injection <input type="checkbox"/> Reclast 5mg injection	<input type="checkbox"/> Xgeva injection <input type="checkbox"/> zoledronic acid 5 mg injection <input type="checkbox"/> Zometa 4 mg injection
Directions: _____	Quantity: _____	Refills: _____	
Diagnosis ( <i>submit documentation</i> ): _____		Dx code ( <i>required</i> ): _____	
1. Does the Recipient have results of a recent bone mineral density test (BMD)?		<input type="checkbox"/> Yes – <i>submit documentation of BMD test results</i> <input type="checkbox"/> No	
2. Based on the US-adapted World Health Organization (WHO) algorithm, does one of the following apply to the Recipient? <input type="checkbox"/> 10-year probability of hip fracture ≥ 3% <input type="checkbox"/> 10-year probability of major fracture related to osteoporosis ≥ 20%		<input type="checkbox"/> Yes – <i>submit all supporting documentation</i> <input type="checkbox"/> No	
3. Was the Recipient evaluated for other possible causes of osteoporosis, including the following laboratory tests? <i>Check all that apply.</i>		<input type="checkbox"/> Yes – <i>submit results of all requested lab tests</i> <input type="checkbox"/> No	
<input type="checkbox"/> CBC <input type="checkbox"/> Vitamin D <input type="checkbox"/> ionized calcium <input type="checkbox"/> phosphorous		<input type="checkbox"/> albumin <input type="checkbox"/> total protein <input type="checkbox"/> creatinine <input type="checkbox"/> liver enzymes/LFTs	<input type="checkbox"/> thyroid stimulating hormone (TSH) <input type="checkbox"/> urinary calcium excretion <input type="checkbox"/> intact parathyroid hormone (PTH) <input type="checkbox"/> testosterone (if male)
4. Does the Recipient have a history of trial and failure, contraindication, or intolerance to the following agents? <i>Check all that apply.</i>		<input type="checkbox"/> Yes – <i>submit all supporting documentation of trial and failure, intolerance, or contraindications</i> <input type="checkbox"/> No	
<input type="checkbox"/> Actonel tablet <input type="checkbox"/> alendronate tablet <input type="checkbox"/> alendronate oral solution <input type="checkbox"/> Atelvia DR tablet <input type="checkbox"/> Binosto tablet		<input type="checkbox"/> Boniva tablet <input type="checkbox"/> Fosamax tablet <input type="checkbox"/> Fosamax Plus D tablet <input type="checkbox"/> ibandronate tablet	<input type="checkbox"/> pamidronate IV <input type="checkbox"/> risedronate tablet <input type="checkbox"/> risedronate DR tablet <input type="checkbox"/> zoledronic acid 4 mg injection
5. <i>For Xgeva requests</i> , does the Recipient have a history of trial and failure, contraindication, or intolerance to the preferred injectable agent, <u>zoledronic acid 4 mg injection</u> ( <i>generic Zometa</i> )?		<input type="checkbox"/> Yes – <i>submit all supporting documentation of trial and failure, intolerance, and contraindications</i> <input type="checkbox"/> No	
6. Injectable Bone Resorption Suppression Agents are part of the Department's Specialty Pharmacy Drug Program (SPDP). What Specialty Pharmacy will be used? Refer to the Department's SPDP website for more information: <a href="http://www.dhs.pa.gov/provider/pharmacyservices/thespecialtypharmacydrugprogram/index.htm">http://www.dhs.pa.gov/provider/pharmacyservices/thespecialtypharmacydrugprogram/index.htm</a> .		<input type="checkbox"/> Diplomat Specialty Pharmacy <input type="checkbox"/> Walgreens Specialty Pharmacy	

**PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION TO DHS – PHARMACY DIVISION**

Prescriber Signature: _____	Date: _____
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