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Please note: All information below is required to process this request.

Mon-Fri: 5am to 10pm Pacific / Sat: 6am to 3pm Pacific

## Forteo® Prior Authorization Request Form (Page 1 of 2)

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Member Information (required)			Provider Information (required)		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:
Medication Information (required)					
Medication Name:			Strength:		Dosage Form:
<input type="checkbox"/> Check if requesting <b>brand</b>			Directions for Use:		
<input type="checkbox"/> Check if request is for <b>continuation of therapy</b>					
Clinical Information (required)					
<p><b>Select the diagnosis below:</b></p> <input type="checkbox"/> Glucocorticoid-induced osteoporosis in men and women at high risk for fracture <input type="checkbox"/> Osteoporosis– To increase bone mass in men with primary or hypogonadal osteoporosis at high risk for fracture <input type="checkbox"/> Osteoporosis– For postmenopausal women at high risk for fracture <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
<p><b>Clinical Information:</b></p> <p>Document the bone mineral density (BMD) T-score from the lumbar spine, femoral neck, total hip, or radius (one-third radius site):            T-Score: _____ (specify if negative) Date: _____</p> <p>Has the treatment duration with parathyroid hormones [Forteo (teriparatide), Tymlos (abaloparatide)] exceeded 24 months of therapy during the patient's lifetime? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p>Please document the number of months of parathyroid hormone therapy the patient has used in his/her lifetime: _____ months</p> <p>Select if the patient has the following FRAX (Fracture Risk Assessment Tool) 10-year probabilities:</p> <input type="checkbox"/> Major osteoporotic fracture at 20% or more in the U.S., or the country-specific threshold in other countries or regions <input type="checkbox"/> Hip fracture at 3% or more in the U.S., or the country-specific threshold in other countries or regions					
<p><b>For glucocorticoid-induced osteoporosis in men and women at high risk for fracture, also answer the following:</b></p> <p>Does the patient have history of taking prednisone, or its equivalent, at a dose <math>\geq</math> 5 mg/day for 3 months or more? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p> <p>Select if the patient has history of fractures resulting from minimal trauma including the following:</p> <input type="checkbox"/> Fracture of the hip <input type="checkbox"/> Fracture of the distal radius <input type="checkbox"/> Fracture of the pelvis <input type="checkbox"/> Fracture of the proximal humerus <input type="checkbox"/> Vertebral compression fracture					
<p>Has the patient had trial and failure, contraindication, or intolerance to one bisphosphonate (e.g., alendronate)? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p>					
<p><b>For osteoporosis in postmenopausal women or men with primary or hypogonadal osteoporosis at high risk for fracture, also answer the following:</b></p> <p>Select if the patient has one of the following diagnosis:</p> <input type="checkbox"/> Postmenopausal osteoporosis or osteopenia <input type="checkbox"/> Primary or hypogonadal osteoporosis or osteopenia <p>Select if the patient has history of fractures resulting from low-trauma including the following:</p> <input type="checkbox"/> Distal forearm <input type="checkbox"/> Hip <input type="checkbox"/> Pelvis <input type="checkbox"/> Proximal humerus <input type="checkbox"/> Spine					
<p>Has the patient had trial and failure, contraindication, or intolerance to one osteoporosis treatment [e.g., alendronate, risedronate, zoledronic acid, Prolia (denosumab)]? <input type="checkbox"/> <b>Yes</b> <input type="checkbox"/> <b>No</b></p>					

This document and others if attached contain information that is privileged, confidential and/or may contain protected health information (PHI). The Provider named above is required to safeguard PHI by applicable law. The information in this document is for the sole use of OptumRx. Proper consent to disclose PHI between these parties has been obtained. If you received this document by mistake, please know that sharing, copying, distributing or using information in this document is against the law. **If you are not the intended recipient, please notify the sender immediately.**  
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**Forteo<sup>®</sup> Prior Authorization Request Form (Page 2 of 2)**  
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Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

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Please note: This request may be denied unless all required information is received.  
For urgent or expedited requests please call 1-800-711-4555.  
This form may be used for non-urgent requests and faxed to 1-800-527-0531.