

Medical prior authorization form

Urgent means the standard review time may seriously jeopardize the life or health of the patient or the patient to regain maximum function. Boniva® (intravenous ibandronate) Member Last Name:	Fax comple This form appli This request is:	_	mercial Individual	· —	
Boniva® (intravenous ibandronate) Member Last Name:					
Member Last Name:		to regain maximum function.	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	,	
Last Name:	Boniva	(intravenous ibandronate)			
D #:	Member				
D #:	Last Name:		First Name:		
Requesting Physician: Phone: Fax:	ID #:				
Physician Address: Physician NPI: Provider Signature: Date: Product and Billing Information Drug product: Boniva 1 mg/mL injection ICD-10 Code(s): Dose:	Primary Care Ph	ysician:	_		
Provider Signature: Product and Billing Information Drug product: Boniva 1 mg/mL injection Dose: bandronate 1 mg/mL injection bandronate 1 mg/ml syringe Date: Date:	Requesting Phys	sician:	Phone:	Fax:	
Product and Billing Information Drug product:					
Product and Billing Information Drug product:	Physician NPI: _				
Drug product: Boniva 1 mg/mL injection ICD-10 Code(s): Ibandronate 1 mg/mL injection Dose: Frequency: Date of last dose: Date of next dose: Date of next dose: New request Continuation request Continuation request Continuation request Start Date: Date of next dose: New request Continuation request Start Date: New request Start Date: Start Date: Start Date: New request Start Date: Start Date:	Provider Signature:		Date:		
Ibandronate 1 mg/mL injection	Product and	Billing Information			
	Drug product:	☐ Boniva 1 mg/mL injection	ICD-10 Code(s):		
Date of last dose: Date of next dose: New request Continuation request Administration: Physician's Office Outpatient Infusion Facility: Home infusion Agency: NPI: Fax #: Billing: Physician Buy and Bill Facility Buy and Bill Specialty Pharmacy		☐ Ibandronate 1 mg/mL injection	Dose:	Frequency:	
Administration: Physician's Office Outpatient Infusion Facility: NPI: Fax #: Home infusion Agency: NPI: Fax #: Billing: Physician Buy and Bill Facility Buy and Bill Specialty Pharmacy		☐ Ibandronate 1 mg/ml syringe	Start Date:		
Administration: Physician's Office Outpatient Infusion Facility: NPI: Fax #: Home infusion Agency: NPI: Fax #: Billing: Physician Buy and Bill Facility Buy and Bill Specialty Pharmacy					
Administration:			Date of next dose:		
□ Outpatient Infusion Facility: NPI: □ Home infusion Agency: NPI: Fax #: Fax #: Billing: Physician Buy and Bill □ Facility Buy and Bill □ Specialty Pharmacy	_		☐ New request ☐ Continuation request		
Facility:	Administration:	-			
☐ Home infusion Agency: NPI: Billing: Physician Buy and Bill ☐ Facility Buy and Bill ☐ Specialty Pharmacy		•	NDI	F. "	
Agency: NPI: Fax #: Billing: Physician Buy and Bill Facility Buy and Bill Specialty Pharmacy			NPI:	Fax #:	
Billing: Physician Buy and Bill Facility Buy and Bill Specialty Pharmacy			NDI.	Fov #1	
☐ Facility Buy and Bill ☐ Specialty Pharmacy	Billing:		INPI	Fax #	
☐ Specialty Pharmacy	Dilling.	-			
			NPI:	Fax #:	
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Before this drug is covered, the patient must meet all of the following criteria:

- 1. Diagnosis of postmenopausal osteoporosis
- 2. Bone mineral density T-score of -2.5 or less
- 3. Must first try alendronate (generic Fosamax), Actonel, ibandronate (generic Boniva tablet), or zoledronic acid (generic Reclast)

For continuation of osteoporosis treatment, patient must meet one of the following requirements:

- **A.** Patient has not received more than 5 years of total treatment with a bisphosphonate or Prolia in a lifetime, unless the patient has a high risk for fracture, such as:
 - Long-term corticosteroid use (7.5 mg prednisone (or equivalent) or higher for 3 months or longer), untreated hypogonadism, spontaneous or surgical premature menopause at less than age 45, hyperparathyroidism, hyperthyroidism, chronic liver disease, patient has epilepsy or is taking anticonvulsant therapy, or a documented fragility fracture
- **B.** Patients without a risk factors present who have received more than 5 years of treatment with a bisphosphonate and/or Prolia, a "drug holiday" is required and:



- a. Patient must be off of osteoporosis treatment for a minimum of two years, and
- b. Patient must have a documented DXA scan showing a significant reduction in bone mineral density (BMD) two years or longer after stopping osteoporosis therapy. Please fax DXA scan results.

Note: Authorization for indications not approved by the Food and Drug Administration (FDA) or recognized in CMS-accepted compendia (e.g. DrugDex, AHFS, U.S. Pharmacopeia, and also Clinical Pharmacology for oncology indications only) require supporting evidence for coverage. Please provide two published peer-reviewed literature articles supporting the drug's use for the identified indication.

Priority Health Precertification Documentation				
Α.	What condition is this drug being used to treat? Postmenopausal osteoporosis Other – the patient's condition is:			
	Rationale for use:			
В.	What other drugs has the patient tried first? alendronate (generic Fosamax) Actonel ibandronate (generic Boniva) zoledronic acid (generic Reclast)			
C.	How many years has the patient's condition been treated with bisphosphonate or Prolia therapy? years, months			
D.	Which of the following, if any, apply to the patient? long-term corticosteroid user taking 7.5 mg of prednisone (or equivalent) or higher for 3 months or longer Untreated hypogonadism Spontaneous or surgical premature menopause at less than age 45 Hyperparathyroidism Hyperthyroidism Chronic liver disease Patient has epilepsy or is taking anticonvulsant therapy Documented fragility fracture Other – the patient's risk factor is:			
If none of the above, has the patient had a two year "drug holiday" from osteoporosis drugs? ☐ Yes Date range of drug holiday:				
	 □ No Does the patient have a documented DXA scan two years or longer after stopping osteoporosis therapy showing a significant reduction in bone mineral density (BMD)? Please fax DXA scan results. □ Yes □ No 			
	What date did the patient stop osteoporosis therapy? Date:			
	What date was the patient's DXA scan was completed on? Date:			