



Miacalcin® (calcitonin) Medication Precertification Request

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(All fields must be completed and legible for Precertification Review.)

Aetna Precertification Notification

Phone: 1-866-752-7021

FAX: 1-888-267-3277

For Medicare Advantage Part B:

Phone: 1-866-503-0857

FAX: 1-844-268-7263

Please indicate: Start of treatment: Start date ____/____/____
 Continuation of therapy: Date of last treatment ____/____/____

Precertification Requested By: _____ Phone: _____ Fax: _____

A. PATIENT INFORMATION

First Name:		Last Name:	
Address:		City:	State: ZIP:
Home Phone:	Work Phone:	Cell Phone:	
DOB:	Allergies:	E-mail:	
Current Weight: _____ lbs or _____ kgs		Height: _____ inches or _____ cms	

B. INSURANCE INFORMATION

Aetna Member ID #: _____	Does patient have other coverage? <input type="checkbox"/> Yes <input type="checkbox"/> No
Group #: _____	If yes, provide ID#: _____ Carrier Name: _____
Insured: _____	Insured: _____
Medicare: <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, provide ID #: _____	Medicaid: <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, provide ID #: _____

C. PRESCRIBER INFORMATION

First Name:		Last Name:		(Check One): <input type="checkbox"/> M.D. <input type="checkbox"/> D.O. <input type="checkbox"/> N.P. <input type="checkbox"/> P.A.	
Address:		City:	State:	ZIP:	
Phone:	Fax:	St Lic #:	NPI #:	DEA #:	UPIN:
Provider E-mail:		Office Contact Name:		Phone:	
Specialty (Check one): <input type="checkbox"/> Gynecologist <input type="checkbox"/> Internal Medicine <input type="checkbox"/> Hematologist <input type="checkbox"/> Endocrinologist <input type="checkbox"/> Other: _____					

D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION

Place of Administration: <input type="checkbox"/> Self-administered <input type="checkbox"/> Physician's Office <input type="checkbox"/> Outpatient Infusion Center Phone: _____ Center Name: _____ <input type="checkbox"/> Home Infusion Center Phone: _____ Agency Name: _____ <input type="checkbox"/> Administration code(s) (CPT): _____ Address: _____	Dispensing Provider/Pharmacy: Patient Selected choice <input type="checkbox"/> Physician's Office <input type="checkbox"/> Retail Pharmacy <input type="checkbox"/> Specialty Pharmacy <input type="checkbox"/> Other: _____ Name: _____ Address: _____ Phone: _____ Fax: _____ TIN: _____ PIN: _____
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E. PRODUCT INFORMATION

Request is for: Miacalcin (calcitonin) Dose: _____ Frequency: _____

F. DIAGNOSIS INFORMATION – Please indicate primary ICD Code and specify any other where applicable.

Primary ICD Code: _____ Secondary ICD Code: _____ Other ICD Code: _____

G. CLINICAL INFORMATION – Required clinical information must be completed in its entirety for all precertification requests.

For Initiation Requests (clinical documentation required):

Hypercalcemia
 Yes No Does the patient have a diagnosis of hypercalcemia?

Management of pain following an osteoporotic vertebral fracture
 Yes No Does the patient have a diagnosis of osteoporosis?
 Yes No Has the osteoporotic spinal compression fracture been verified on imaging with the correlating clinical signs and symptoms suggesting an acute injury?
→ Please indicate date of event or onset of symptoms: ____/____/____
Please indicate date of imaging: ____/____/____ (Imaging report must be submitted for review)

Yes No Has the patient had an ineffective response, intolerance, or contraindication to standard analgesic therapy (e.g., non-steroidal anti-inflammatory drug (NSAID), acetaminophen)?
 Yes No Has the patient had an intolerance or contraindication to intranasal calcitonin?
 Yes No Unknown Is the patient neurologically intact?

Paget's disease of bone
 Yes No Does the patient have symptoms of Paget's disease of bone (e.g., bone pain, bone deformity, fracture, hearing loss)?
 Yes No Has the patient failed prior treatment with an injectable bisphosphonate (e.g., pamidronate, Reclast (zoledronic acid)) or experienced intolerance to a previous injectable bisphosphonate?

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Patient First Name	Patient Last Name	Patient Phone	Patient DOB
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G. CLINICAL INFORMATION (continued) – Required clinical information must be completed in its entirety for all precertification requests.

Postmenopausal osteoporosis

- Yes No Is the patient greater than 5 years post-menopause?
- Yes No Does the patient have a history of fragility fractures?
 Please indicate the patient's pre-treatment T-score:
 -2.5 or below (e.g., -2.6, -2.7, -3) Osteopenia [between -2.5 and -1 (e.g., -2.4, -2.3, -2)] -1 or above (e.g., -0.9, -0.8, -0.5) unknown
 Please indicate the patient's pre-treatment FRAX score for any major fracture: less than 20% greater than or equal to 20% unknown
 Please indicate the patient's pre-treatment FRAX score for hip fracture: less than 3% greater than or equal to 3% unknown
- Yes No Has the patient failed prior treatment of an oral bisphosphonate?
 Yes No Is there a clinical reason to avoid treatment with a bisphosphonate?
 Please indicate clinical reason:
 presence of anatomic or functional esophageal abnormalities that might delay transit of the tablet (e.g., achalasia, stricture, or dysmotility)
 active upper gastrointestinal problem (e.g., dysphagia, gastritis, duodenitis, erosive esophagitis, ulcers)
 presence of documented or potential gastrointestinal malabsorption (e.g., gastric bypass procedures, celiac disease, Crohn's disease, infiltrative disorders, etc.)
 inability to stand or sit upright for 30 to 60 minutes
 inability to take oral bisphosphonate at least 30 to 60 minutes before first food, drink or medication of the day
 renal insufficiency (creatinine clearance less than 35 ml/min)
 history of intolerance to an oral bisphosphonate
 no clinical reason to avoid treatment with an oral bisphosphonate
 other, please explain: _____
- Yes No Has the patient failed prior treatment with or is intolerant to previous injectable osteoporosis therapy (e.g., zoledronic acid [Reclast], teriparatide [Forteo, Bonsity], denosumab [Prolia], abaloparatide [Tymlos])?
- Yes No Has the patient had an intolerance or contraindication to intranasal calcitonin?

For Continuation Requests (clinical documentation required):

Hypercalcemia

- Yes No Has the patient experienced benefit from therapy as evidenced by disease stability or disease improvement?
- Yes No Has the patient experienced any adverse effects?

Management of pain following an osteoporotic vertebral fracture

- Yes No Does the patient have a diagnosis of osteoporosis?
- Yes No Has the osteoporotic spinal compression fracture been verified on imaging with the correlating clinical signs and symptoms suggesting an acute injury?
 Please indicate date of event or onset of symptoms: ____/____/____
 Please indicate date of imaging: ____/____/____ (Imaging report must be submitted for review)
- Yes No Has the patient had an ineffective response, intolerance, or contraindication to standard analgesic therapy (e.g., non-steroidal anti-inflammatory drug (NSAID), acetaminophen)?
- Yes No Has the patient had an intolerance or contraindication to intranasal calcitonin?
- Yes No Unknown Is the patient neurologically intact?

Paget's disease of bone

- Yes No Is the patient currently receiving the requested medication through samples or a manufacturer's patient assistance program?
- Yes No Has the patient experienced benefit from therapy as evidence by disease stability or disease improvement?
- Yes No Has the patient experienced any adverse effects?

Postmenopausal osteoporosis

- Yes No Is the patient currently receiving the requested medication through samples or a manufacturer's patient assistance program?
- Please indicate the length of time on therapy:
 Less than 24 months
 Yes No Is the patient experiencing clinical benefit as evidenced by no adverse events during therapy (i.e., no clinically significant adverse reaction, no new fracture seen on radiography)?
- 24 months or greater
 Yes No Has the patient experienced clinical benefit as evidenced by a bone mass measurement showing an improvement or stabilization in T-score compared with the previous bone mass measurement?
 Yes No Has the patient experienced any adverse effects?

H. ACKNOWLEDGEMENT

Request Completed By (Signature Required): _____ Date: ____/____/____

Any person who knowingly files a request for authorization of coverage of a medical procedure or service with the intent to injure, defraud or deceive any insurance company by providing materially false information or conceals material information for the purpose of misleading, commits a fraudulent insurance act, which is a crime and subjects such person to criminal and civil penalties.

The plan may request additional information or clarification, if needed, to evaluate requests.