



# Tymlos™ (abaloparatide) Medication Precertification Request

Aetna Precertification Notification  
Phone: 1-855-240-0535  
FAX: 1-877-269-9916

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

For Medicare Advantage Part B:  
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:	Email:	
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 #:
Provider Email:	Office Contact Name:	Phone:		
Specialty (Check one): <input type="checkbox"/> GYN <input type="checkbox"/> Orthopedic <input type="checkbox"/> Primary Provider <input type="checkbox"/> Other: _____				

### D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION

<b>Place of Administration:</b> <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: _____	<b>Dispensing Provider/Pharmacy: Patient Selected choice</b> <input type="checkbox"/> Physician's Office <input type="checkbox"/> Retail Pharmacy <input type="checkbox"/> Specialty Pharmacy <input type="checkbox"/> Mail Order <input type="checkbox"/> Other: _____ Name: _____ Address: _____ Phone: _____ Fax: _____ TIN: _____ PIN: _____
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### E. PRODUCT INFORMATION

Request is for Tymlos: 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 All Requests (clinical documentation required)**  
Please provide the patient's Bone Mineral Density (BMD) score and date obtained: T-score: \_\_\_\_\_ Date: \_\_\_\_ / \_\_\_\_ / \_\_\_\_

Yes  No Does the patient have clinical evidence of uncorrected preexisting hypocalcemia?  
 Yes  No Is the patient at high risk for fractures?  
 Yes  No Has the cumulative use of Tymlos (abaloparatide) and parathyroid hormone analogs (e.g., teriparatide) exceeded 2 years duration?

**For initiation requests:**  
**Post-menopausal osteoporosis**  
 Yes  No Has the patient had an osteoporotic fracture?  
 Yes  No Does the patient have multiple risk factors for fractures?  
→ Please select all that apply:  anorexia nervosa  alcohol intake of 4 or more units a day  corticosteroid therapy  smoking  
 Cushing's syndrome  failed previous osteoporosis therapy  high risk for falls  history of osteoporosis fractures  
 increasing age  intolerant to previous osteoporosis therapy  low body mass  parental history of hip fracture  
 rheumatoid arthritis  other: please explain: \_\_\_\_\_

Yes  No Is there clinical evidence that a trial of two bisphosphonates was ineffective?  
→  Yes  No Is there clinical evidence that a trial of at least one bisphosphonate and one selective estrogen receptor modulator (SERM)?  
→ Please identify the failure of the medication trial:  Continued bone loss  Other: please identify: \_\_\_\_\_

→ Please select which of the following bisphosphonates and/or SERM's the patient tried  
**Select all that apply:**  Actonel or Actonel with Calcium (risedronate)  Atelvia (risedronate)  Boniva (ibandronate)  
 Didronel (etidronate)  Fosamax or Fosamax plus D (alendronate)  Skelid (tiludronate)  
 Zometa/Reclast (zoledronic acid)  Fareston (toremifene)  Tamoxifen (nolvadex)  
 Evista (raloxifene)  Other: Please identify: \_\_\_\_\_

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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 for ALL precertification requests.**

Bisphosphonate #1 Date range: \_\_\_\_/\_\_\_\_/\_\_\_\_ to \_\_\_\_/\_\_\_\_/\_\_\_\_

Bisphosphonate #2 OR SERM Date range: \_\_\_\_/\_\_\_\_/\_\_\_\_ to \_\_\_\_/\_\_\_\_/\_\_\_\_

Yes  No Is there documented evidence that the patient has an intolerance to bisphosphonates and/or SERMs?

→ **Select all that apply: Bisphosphonates:**  Persistent upper GI disturbance  Severe musculoskeletal pain  Hypocalcemia  
 Other: please explain: \_\_\_\_\_

**SERM:**  Flu Syndrome  Hot flashes  Nausea/vomiting or diarrhea  Arthralgia  Rhinitis  
 Other: please specify: \_\_\_\_\_

Yes  No Is there documented evidence that the patient has a contraindication to bisphosphonates and/or SERMs?

→ **Select all that apply: Bisphosphonates:**  Renal Impairment  Hypersensitivity to bisphosphonates or components  
 Other: please identify: \_\_\_\_\_

**SERM:**  Active or history of venous thromboembolism (e.g. DVT, PE, RVT)  Hypersensitivity  
 Hx. of CVA or TIA  Other: please identify: \_\_\_\_\_

**For continuation requests:**

Yes  No Does the patient have a hypersensitivity to Tymlos (abaloparatide)?

→ Please indicate which of the following reactions the patient had:

anaphylaxis  dyspnea  facial and upper airway edema  hypotension  pruritus  rash  urticaria  
 other: \_\_\_\_\_

Please indicate what type of response the patient has experienced on therapy:

**Select appropriate response:**  No response  Minimal response  Adequate response  Significant improvement

**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.