

## ALENDRONATE / RALOXIFENE / RISEDRONATE FOR OSTEOPOROSIS SPECIAL AUTHORIZATION REQUEST FORM

Please complete all required sections to allow your request to be processed.

Patients may or may not meet eligibility requirements as established by Alberta Government sponsored drug programs.

PATIENT INFORMATION						COVERAGE TYPE:	
PATIENT LAST NAME	FIRST NAME				INITIAL	☐ Alberta Blue Cross ☐ Alberta Human Services	
DATE OF BIRTH: Year / Month / Day	ALBERTA PERSONAL HEALT			LTH NUMBER		Other	
STREET ADDRESS	ADDRESS CITY			PROV	POSTAL CODE	IDENTIFICATION/CLIENT/COVERAGE No:	
PRESCRIBER INFORMATION							
PRESCRIBER LAST NAME FIRST NAME INITIAL				PRESCRIBER PROFESSIONAL ASSOCIATION REGISTRATION			
				☐ CPSA ☐ ACO REGISTRATION NO.			
STREET ADDRESS				☐ CARNA ☐ ADA+C ☐ ACP ☐ Other			
CINEEL NOONES				ONE:	☐ Other	FAV.	
CITY, PROVINCE				JINE.		FAX:	
POSTAL CODE				FAX NUMBER MUST BE PROVIDED WITH EACH REQUEST SUBMITTED			
Criteria for Coverage							
"For the treatment of osteoporosis in patients with a 20% or greater 10-year fracture risk who have documented intolerance to alendronate 70 mg or risedronate 35 mg. Special authorization may be granted for 6 months."							
"Requests for other osteoporosis medications covered via special authorization will not be considered until 6 months after the last dose of denosumab 60 mg/syr injection syringe."							
"Requests for other osteoporosis medications covered via special authorization will not be considered until 12 months after the last dose of zoledronic acid 0.05 mg/ml injection."							
Note: The fracture risk can be determined by the World Health Organization's fracture risk assessment tool, FRAX, or the most recent (2010) version of the Canadian Association of Radiologist and Osteoporosis Canada (CAROC) table.							
* Alendronate 70 mg and risedronate 35 mg are regular benefits not requiring Special Authorization.							
** Alendronate and risedronate also have special authorization criteria for Paget's disease. Please refer to the Alberta Drug Benefit List for							
alendronate and risedronate's other criteria for the indication of Paget's disease: <a href="http://www.health.alberta.ca/services/drug-benefit-list.html">http://www.health.alberta.ca/services/drug-benefit-list.html</a> Please provide the following information for ALL requests:							
Indicate which drug is requested (check ONE box): Alendron					Raloxifene	Risedronate	
Please provide the following information for all NEW requests:							
Diagnosis: ☐ For the treatment of Osteoporosis ☐ Osteopenia ☐ Other, please specify:							
Fracture risk:							
a) Has the patient experienced FRACTURES related to the diagnosis?							
b) Does the patient have a 20% or greater 10-year fracture risk?  No Yes							
Information regarding previous alendronate 70mg or risedronate 35mg use:							
alendronate 70mg or risedronate 35mg HAS been utilized.							
Nature of response:							
Other (please specify):							
alendronate 70mg or risedronate 35mg has NOT been utilized (specify reason(s)):							
RENEWAL: This product is eligible for auto-renewal for treatment of osteoporosis. A Special Authorization renewal request is required only if the Special Authorization approval for treatment of osteoporosis has lapsed (i.e. the patient has <u>not</u> made a claim for the drug product during the Approval Period).							
• A				forward this request to: Iberta Blue Cross, Clinical Drug Services 0009-108 Street NW, Edmonton, Alberta T5J 3C5			
		•				1-877-828-4106 toll-free all other areas	
ONCE YOUR REQUEST HAS SUCCESSFULLY TRANSMITTED, PLEASE DO NOT MAIL OR RE-FAX YOUR REQUEST.							