

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	<input type="checkbox"/> Alberta Blue Cross <input type="checkbox"/> Alberta Human Services <input type="checkbox"/> Other	
DATE OF BIRTH: Year / Month / Day	ALBERTA PERSONAL HEALTH NUMBER			
STREET ADDRESS	CITY	PROV	POSTAL CODE	IDENTIFICATION/CLIENT/COVERAGE No:

PRESCRIBER INFORMATION			
PRESCRIBER LAST NAME	FIRST NAME	INITIAL	PRESCRIBER PROFESSIONAL ASSOCIATION REGISTRATION
			<input type="checkbox"/> CPSA <input type="checkbox"/> ACO REGISTRATION NO. <input type="checkbox"/> CARNA <input type="checkbox"/> ADA+C <input type="checkbox"/> ACP <input type="checkbox"/> Other
STREET ADDRESS			PHONE:
CITY, PROVINCE			FAX:
POSTAL CODE			FAX NUMBER MUST BE PROVIDED WITH EACH REQUEST SUBMITTED

Indicate which drug is requested (check ONE box): Denosumab 60 mg/syr Zoledronic Acid 0.05 mg/ml

Indicate diagnosis and associated risk factors:

<input type="checkbox"/> POSTMENOPAUSAL OSTEOPOROSIS → Please indicate which of the following pertain to this patient (check ALL that apply): <input type="checkbox"/> prior fragility fracture <input type="checkbox"/> bone mineral density (BMD) T-Score of less than or equal to -2.5	<input type="checkbox"/> MALE OSTEOPOROSIS → Please indicate which of the following pertain to this patient (check ALL that apply): <input type="checkbox"/> high 10-year risk (i.e., greater than 20%) of experiencing a major osteoporotic fracture <input type="checkbox"/> moderate 10-year fracture risk (10-20%) <input type="checkbox"/> prior fragility fracture
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Other (specify):

Please indicate which of the following pertain to this patient (check ALL that apply):

Denosumab requests only

- bisphosphonates are contraindicated due to drug-induced hypersensitivity (i.e., immunologically mediated).
- bisphosphonates are contraindicated due to severe renal impairment (i.e. creatinine clearance < 35 mL/min)

Denosumab and Zoledronic Acid requests

- abnormality of the esophagus which delays esophageal emptying
- severe gastrointestinal intolerance* with previous use of the following bisphosphonates. If so, must specify the nature and severity of adverse effects:
 alendronate: _____
 risedronate: _____

*Note: severe gastrointestinal intolerance is defined as manifested by weight loss or vomiting directly attributable to the oral bisphosphonate

- unsatisfactory response (defined as a fragility fracture despite adhering to oral alendronate or risedronate treatment fully for 1 year and evidence of a decline in BMD below pre-treatment baseline level)

Additional information relating to request:

PRESCRIBER'S SIGNATURE	DATE	Please forward this request to: ▪ Alberta Blue Cross, Clinical Drug Services 10009-108 Street NW, Edmonton, Alberta T5J 3C5 ▪ FAX: 780-498-8384 in Edmonton • 1-877-828-4106 toll-free all other areas
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ONCE YOUR REQUEST HAS SUCCESSFULLY TRANSMITTED, PLEASE DO NOT MAIL OR RE-FAX YOUR REQUEST.

