

Rank Ligand (RANKL) Inhibitors

Please provide the information below, please print your answer, attach supporting documentation, sign, date, and return to our office as soon as possible.

Please FAX responses to: (800) 869-7791. Phone: (855) 322-4082

Date of request:					
Patient Date of birth			Molina ID		
Pharmacy name	Pharmacy NPI	Telephone number		Fax number	
Prescriber	Prescriber NPI	Telep	hone number	Fax number	
Medication and strength		Directions for use		Qty/Days supply	
1. Is this request for a continuation of existing therapy? Yes No If yes, is there documentation demonstrating disease stability or a positive clinical response)? Yes No 2. Indicate patient's diagnosis: Glucocorticoid-induced osteoporosis Postmenopausal osteoporosis Bone loss in men with prostate cancer Bone loss in women with breast cancer Bone metastasis from solid tumors Multiple myeloma with skeletal-related events Giant cell tumor of bone Hypercalcemia of malignancy					
 3. Will the medication be used in combination with other bone density regulators? Yes No If yes, specify: bisphosphonates raloxifene Prolia (denosumab) Xgeva (denosumab) 					
 Indicate if patient has any of the following: □ Presence of fragility fractures of the hip or spine regardless of bone mineral density □ T-score ≤ -2.5 in the lumbar spine, femoral neck, total hip □ T-score between -1 and -2.5 with a history of recent fragility fracture of proximal humerus, pelvis, or distal forearm 					

fra	T-score between -1 and -2.5 with a FRAX 10-year probability for major fracture ≥20% or hip acture ≥3%
unl	s the patient been treated with at least one Apple Health Preferred Drug (oral or intravenous) ess ineffective, contraindicated or not tolerated? Please select all that apply: Bisphosphonate (minimum trial of 12 months), specify: Selective estrogen receptor modulator (SERM) (minimum trial of 24 months), specify:
	Other, specify: Contraindicated, provide contraindication:
6. Wil equ <u>If</u> y	iagnosis of Glucocorticoid Induced Osteoporosis: Il patient be initiating or continuing systemic glucocorticoid therapy at a daily dosage uivalent to ≥ 7.5 mg of prednisone? ☐ Yes ☐ No res, is patient expected to remain on glucocorticoid therapy for at least 6 months? Yes ☐ No
7. Is prelu	loss in men and prostate cancer: catient currently receiving androgen deprivation therapy (ADT) (e.g., leuprolide, degarelix, agolix) for non-metastatic prostate cancer? Yes No Contraindicated or not tolerated. Explain:
8. Wil letr 	loss in women with breast cancer: Il patient be receiving adjuvant aromatase inhibitor therapy (e.g., anastrozole, exemestane, rozole) for breast cancer? Yes No Contraindicated or not tolerated. Explain:
9. Do	ple Myeloma: es patient have a history of failure, contraindication, or intolerance to zoledronic acid? Yes
10. Ind	cell tumor of bone: licate the following for patient. Check all that apply. Disease is unresectable or surgical resection is likely to result in severe morbidity? Disease recurrent or metastatic
	rcalcemia of malignancy es patient have a baseline corrected serum calcium > 12.5 mg/dL? Yes No

MHW Part#0062RX-2505

MHW-05/02/2025, HCA-02/19/2025 (30.04.45)

CHART NOTES ARE REQUIRED WITH THIS REQUEST					
Prescriber signature	Prescriber specialty	Date			