

Xgeva Prior Authorization Form

Incomplete forms will not be reviewed

Maryland Medicaid Pharmacy Program

Fax: (866) 440-9345 Phone: (800) 932-3918

			Date (MM/DD/YYYY):			
Patient	t Infor	rmation				
Name:			DOB (MM/DD/YYYY):			
		sistance Number:nformation	Sex: ☐ M	☐ F	Height:	Weight:
Name:			NPI:			
Phone:		Fax:				
		son for this Request:				
Name: _		Phone:			Fax:	
Dose:	geva®	120 mg subcutaneously every 4 weeks, with additional	doses on days 8 a	and 15 c	of the first mo	nth of treatment.
☐ Xgeva®120 mg subcutaneously every 4 weeks.						
Yes	No	X	Xgeva [®]			
		Does the patient have a diagnosis of giant cell tumor of the bone?				
		Does the patient have a tumor that is either recurrent, unresectable or where surgical resection is likely to cause severe morbidity?				
		Is the patient a skeletally mature adolescent with a weight > 45 kg?				
		Does that patient have a diagnosis of bone metastases from solid tumors?				
		Does the patient have a diagnosis of hypercalcemia of malignancy?				
		Is the hypercalcemia refractory to intravenous bisphosphonate therapy?				
		Does the patient have any contraindications to Xgeva® therapy (hypocalcemia, hypersensitivity to ingredients)?				
		Is patient currently receiving Prolia [®] ? Xgeva [®] includes the same active ingredient (denosumab) found in Prolia [®] . Patients receiving Xgeva [®] should not take Prolia [®] .				
I attest	that:					
accurat	e to th	its of the treatment outweigh the risks and verify that ne best of my knowledge and prescriber acknowledge and agree that this reque		_		
		d as an original signature for all purposes and shall ha	•	•	•	
D.,, '1	ء	2:	Data		١.	
rrescri	Der S S	Signature:	Date (MM/DD/YYYY):			