

## PROLIA **(FEMALE)** AUTHORIZATION AND RE-AUTHORIZATION REQUEST

TEL: (716) 929-1000 | 1-800-809-4763 FAX: (716) 532-7360

Member Name: Todav's D		Today's Date:	: Date Needed:	
		,		
Date of birth: Sex:	Weight:		Prescriber:	Specialty:
Home Phone Number:			Phone Number:	Fax Number:
( )			( )	( )
Home Address:	City: State	e: Zip:	Address:	City: State: Zip:
☐ Independent Health ☐ Anne Arundel Health System ☐ Pharmacy Benefit Dimensions	<ul><li>☐ Commercial</li><li>☐ Medicaid</li></ul>	<ul><li>☐ Medicare</li><li>☐ Self-funded</li></ul>	Allergies:	Medication ships to patient home
Insurance ID:	Group Number:			Medication ships to provider office
	ST	ATEMENT OF M	EDICAL NECESSITY	
☐ New Authorization	☐ Re-authorization	on*	Dose:	Frequency:
Primary Diagnosis:  Date of Diagnosis:  ICD10 Code:  Prior Treatments:			ip <b>DR</b> □ Patient has a history of <b>NND</b>	an or equal to -1.0 at either the lumbar spine or total osteoporotic fracture calcium level which is within normal limits is
Is the patient female?			AND  □ Documentation showing that patient has been instructed about the symptoms of hypocalcemia and the importance of adequate calcium and vitamin D supplementation while on this therapy is submitted  AND  Patient has demonstrated at least one of the below:  □ Tried and failed oral alendronate therapy as evidenced by disease progression  OR  □ Has documented inability to swallow or established esophageal	
AND			diagnosis which prevents oral administration of alendronate  OR  □ Female patient receiving concurrent adjuvant AI therapy for hormone receptor positive breast cancer.	
<ul><li>□ Patient is at high risk of fracture, defined as</li><li>□ History of osteoporotic fracture <b>OR</b></li></ul>				
Multiple risk factors for fracture (at least two of the following):  Limited movement, such as using wheelchair.  History of frequent falls  Medical condition likely to cause bone loss:			AND  ☐ The patient is at high risk for fracture across multiple skeletal sites, having a BMD T-score at the lumbar spine, total hip, or femoral neck of less than -1.0 or a history of osteoporotic fracture	
□ Concurrent use of medications that may cause bone loss:     □ Concurrent use of medications that may increase risk of falls:			AND  ☐ Patient's current serum calcium level which is within normal limits is submitted	
OR  ☐ Patient is receiving treatment for glucocorticoid-induced osteoporosis  AND			AND  ☐ Documentation showing that patient has been instructed about the symptoms of hypocalcemia and the importance of adequate calcium and vitamin D supplementation while on this therapy is submitted	
☐ Patient will be initiating or continuing systemic glucocorticoid therapy at a daily dosage equivalent to greater than or equal to 7.5 mg of prednisone and is expected to remain on glucocorticoid therapy for at least 6 months.			For Re-Authorization:  Submission of BMD-T Score (bi-annually) Date:	
AND			Submission of Serum Ca+ level (annually) Date:	
			For patients who requested Prolia as a treatment for increased bone mass in prostate/breast cancer, continued ADT or AI therapy?	