

## Denosumab (Prolia) Osteoporosis Order Set

 Phone (616) 394-3547  
 Fax (616) 394-2139

 = Optional Order     = Routine Order    (Cross out and initial **BULLETED ORDERS** that do not apply)

### ORDERS

Date: \_\_\_\_\_ Time: \_\_\_\_\_

Patient Name: \_\_\_\_\_ Date of Birth: \_\_\_\_\_ Weight: \_\_\_\_\_ (kg)

Allergies: \_\_\_\_\_

Exclusion Criteria: hypersensitivity (systemic) to denosumab or any component of the formulation; preexisting hypocalcemia; pregnancy.

Diagnosis Code: (ICD-10) \_\_\_\_\_

- Diagnosis:** Treatment of postmenopausal women with osteoporosis at high risk for fracture.
- Treatment to increase bone mass in men with osteoporosis at high risk for fracture.
- Treatment of glucocorticoid-induced osteoporosis in men and women at high risk for fracture.
- Treatment to increase bone mass in men at high risk for fracture receiving androgen deprivation therapy for nonmetastatic prostate cancer.
- Treatment to increase bone mass in women at high risk for fracture receiving adjuvant aromatase inhibitor therapy for breast cancer.

- Administer denosumab (Prolia) 60mg subcutaneously times one.

### PRE-INJECTION / LAB ORDERS DIAGNOSTICS & TESTS

#### Results

*(Within 45 days of administering Prolia)*

DATE	SERUM CREATININE	DATE	SERUM CALCIUM (NORMAL 8.6 – 10.6 MG/DL)	CALCIUM WDL? **
				<input type="checkbox"/> Yes

\*\*Hold medication if serum calcium level is not within normal range.

### NURSING MONITORING PARAMETERS

- Administer denosumab (Prolia) by subcutaneous technique in the upper lateral arm, abdomen, or lateral thigh. See package insert for proper subcutaneous administration and site selection.
- Do not administer denosumab (Prolia) intraarterially, intra-muscularly or intravenously.
- Counsel the patients to take calcium 1200mg daily with at least 800 units of vitamin D daily

- By signing this order, the Medication Guide and Patient Counseling Chart per FDA REMS Requirement for Prolia has been reviewed with the patient and handed to her/him including the serious risks of Prolia and symptoms of each risk.**

Provider Signature: \_\_\_\_\_ Date: \_\_\_\_\_

Provider Name and Credentials (please print): \_\_\_\_\_ Time: \_\_\_\_\_

Office Phone: \_\_\_\_\_ Office Fax: \_\_\_\_\_

