

**Denosumab/Denosumab-bbdz/ Denosumab-bmwo (Prolia/Jubbonti/Stobolco)**  
**Osteoporosis Order Set**☐ = 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.

**MEDICATIONS**

- ☐ Denosumab (Prolia) 60mg subcutaneously times one.
- ☐ Denosumab-bbdz (Jubbonti) 60mg subcutaneously times one.
- ☐ Denosumab-bmwo (Stoboclo) 60mg subcutaneously times one.

**PRE-INJECTION / LAB ORDERS****DIAGNOSTICS & TESTS****Results***(Within 45 days of administering Prolia/Jubbonti)*

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/denosumab-bbdz/denosumab-bmwo (Prolia/Jubbonti/Stoboclo) 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/denosumab-bbdz/denosumab-bmwo (Prolia/Jubbonti/Stoboclo) intraarterially, intramuscularly 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/Jubbonti/Stoboclo has been reviewed with the patient and handed to her/him including the serious risks of Prolia/Jubbonti/Stoboclo and symptoms of each risk.

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

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

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

