



Fax your completed order to 877-734-1157

Patient Name: \_\_\_\_\_ DOB: \_\_\_\_/\_\_\_\_/\_\_\_\_  
Date of Last Infusion: \_\_\_\_/\_\_\_\_/\_\_\_\_ Height \_\_\_\_\_ Weight \_\_\_\_\_  
Infusion Location: (state and site) \_\_\_\_\_

### Evenity (romosozumab-aqqg) Orders

Diagnosis (please provide ICD-10 code in space provided):	
_____ Post-menopausal osteoporosis (ICD-10)	_____ Other: _____ (ICD-10)
<input type="checkbox"/> Provider: Please confirm patient has NOT had an MI or stroke in the past year by checking box	
Recent calcium level: _____ mg/dl Date of result: _____ (please include copy)	

#### Nursing Orders

-Hold treatment and notify provider for:

- Hold for hypocalcemia at initiation of treatment
- Ensure patient is taking daily calcium and Vitamin D supplement
- Planned/recent invasive dental procedures
- Jaw, thigh or groin pain, or dermatologic changes since starting Evenity
- A history of severe bone, muscle or joint pain following Evenity injections

Evenity 210mg subcutaneously in the upper arm, abdomen or upper thigh.

Provided as 2 separate 105mg/1.17ml prefilled syringes. Rotate site with each injection.

#### Frequency:

- Repeat once a month for 12 months
- Other: \_\_\_\_\_

#### Observation Period:

- Following initial Evenity injection, observe patient for 15 minutes for hypersensitivity. Patients who have previously received and tolerated Evenity do not require observation period.
- If reaction occurs, initiate Hypersensitivity Reaction Management Policy/Protocol as clinically indicated.

Provider Name (print) \_\_\_\_\_ Date: \_\_\_\_\_

Provider Signature: \_\_\_\_\_ Time: \_\_\_\_\_