
 <p style="text-align: center;"><b>Oregon Health &amp; Science University Hospital and Clinics Provider's Orders</b></p> <div style="display: flex; align-items: center;"> <div style="writing-mode: vertical-rl; transform: rotate(180deg); font-size: small; margin-right: 5px;">PO7071</div>  </div> <p style="text-align: center;"><b>ADULT AMBULATORY INFUSION ORDER Romosozumab-aqqg (EVENTITY) Injection</b></p> <p style="text-align: center; font-size: small;">Page 1 of 3</p>	<p>ACCOUNT NO. _____</p> <p>MED. REC. NO. _____</p> <p>NAME _____</p> <p>BIRTHDATE _____</p> <p style="text-align: right; font-size: small;"><i>Patient Identification</i></p>
<b>ALL ORDERS MUST BE MARKED IN INK WITH A CHECKMARK ( ✓ ) TO BE ACTIVE.</b>	

**Weight:** \_\_\_\_\_ kg      **Height:** \_\_\_\_\_ cm

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

**Diagnosis Code:** \_\_\_\_\_

**Treatment Start Date:** \_\_\_\_\_      **Patient to follow up with provider on date:** \_\_\_\_\_

**\*\*This plan will expire after 365 days at which time a new order will need to be placed\*\***

**GUIDELINES FOR ORDERING**

1. Send **FACE SHEET and H&P or most recent chart note.**
2. Romosozumab may increase the risk of MI, stroke, and cardiovascular death. It should not be initiated in patients who have had a myocardial infarction or stroke within the preceding year. Consider whether the benefits outweigh the risks in patients with other cardiovascular risk factors.
3. Duration of therapy is limited to 12 monthly doses.
4. Confirm patient has had recent oral/dental evaluation if indicated prior to initiating therapy.
5. Hypocalcemia must be corrected prior to initiation of therapy. All patients should be prescribed daily calcium and Vitamin D supplementation.
6. Risk versus benefit regarding osteonecrosis of the jaw and hip fracture must be discussed prior to treatment.
7. A complete metabolic panel is recommended and a calcium level must be obtained within 30 days prior to starting treatment.
8. **Provider must review and check box for the following question:** Osteonecrosis of the jaw (ONJ) has been reported in patients receiving this medication. Risk factors include higher doses, duration of therapy greater than 2 years, cancer diagnosis, concurrent immunosuppression, poor oral hygiene, periodontal disease, ill-fitting dentures, and invasive dental procedures (among others). Provider attests that patient has had a dental evaluation or has no contraindications to therapy related to dental issues prior to initiating therapy.
  - Osteonecrosis of the jaw (ONJ) risk reviewed prior to initiation of therapy?

**LABS:**

- Complete Metabolic Panel, Routine, ONCE, every 8 weeks.

**NURSING ORDERS:**

1. TREATMENT PARAMETER #1 – When labs are ordered:
  - a. If serum calcium is greater than or equal to 8.4 mg/dL: OK to proceed with treatment.
  - b. If serum calcium less than 8.4 mg/dL AND albumin is less than 4.35 mg/dL: Calculate corrected calcium: Hold treatment and notify provider if corrected calcium is less than 8.4 mg/dL.
  - c. If serum calcium less than 8.4 mg/dL AND albumin is greater than or equal to 4.35 mg/dL: Hold treatment and notify provider.
2. TREATMENT PARAMETER #2 - Assess for new or unusual thigh, hip, groin, or jaw pain. Hold treatment and Contact provider if positive findings. Invasive dental work includes dentoalveolar procedures such as tooth extraction, root canal, or placement of dental implants. Hold treatment and notify provider if the following has not been discussed with the ordering provider:



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- a. If patient is anticipating invasive dental work in the next 3 months
- b. Has received invasive dental work in the last 8 weeks
- c. Has a new diagnosis of severe periodontal disease
3. Please remind patient to take at least 500 mg elemental calcium twice daily and 400 IU Vitamin D daily.
4. RN to assess for previous myocardial infarction (MI) or stroke at every visit. Hold and contact provider if patient had a MI or stroke. Romosozumab-aqqg may increase the risk of MI, stroke, and cardiovascular death. If a patient experiences a MI or stroke during therapy, romosozumab-aqqg should be discontinued.
5. Follow facility policies and/or protocols for vascular access maintenance with appropriate flush solution, declotting (alteplase), and/or dressing changes.

**MEDICATIONS:**

- Romosozumab-aqqg (EVENTITY) 210 mg injection, subcutaneous, ONCE, every 4 weeks for 12 doses

Allow syringes to sit at room temperature for at least 30 minutes before use. Inject two 105 mg/1.17 mL syringes for a total dose of 210 mg. Administer into the thigh, abdomen (except for a 2 inch area around the navel), or outer area of upper arm. Rotate injection sites.

**HYPERSENSITIVITY MEDICATIONS:**

1. NURSING COMMUNICATION – If hypersensitivity or infusion reactions develop, temporarily hold the infusion and notify provider immediately. Administer emergency medications per the Treatment Algorithm for Acute Infusion Reaction (OHSU HC-PAT-133-GUD, HMC C-132). Refer to algorithm for symptom monitoring and continuously assess as grade of severity may progress.
2. diphenhydrAMINE (BENADRYL) injection, 25-50 mg, intravenous, AS NEEDED x 1 dose for hypersensitivity or infusion reaction
3. EPINEPHrine HCl (ADRENALIN) injection, 0.5 mg, intramuscular, AS NEEDED x 1 dose for hypersensitivity or infusion reaction
4. hydrocortisone sodium succinate (SOLU-CORTEF) injection, 100 mg, intravenous, AS NEEDED x 1 dose for hypersensitivity or infusion reaction
5. famotidine (PEPCID) injection, 20 mg, intravenous, AS NEEDED x 1 dose for hypersensitivity or infusion reaction

**By signing below, I represent the following:**

I am responsible for the care of the patient (*who is identified at the top of this form*);

I hold an active, unrestricted license to practice medicine in:  Oregon  \_\_\_\_\_ (*check box that corresponds with state where you provide care to patient and where you are currently licensed. Specify state if not Oregon*);

**My physician license Number is # \_\_\_\_\_ (MUST BE COMPLETED TO BE A VALID PRESCRIPTION);** and I am acting within my scope of practice and authorized by law to order Infusion of the medication described above for the patient identified on this form.

**Provider signature:** \_\_\_\_\_ **Date/Time:** \_\_\_\_\_

**Printed Name:** \_\_\_\_\_ **Phone:** \_\_\_\_\_ **Fax:** \_\_\_\_\_



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Central Intake:

Phone: 971-262-9645 (providers only) Fax: 503-346-8058

***Please check the appropriate box for the patient's preferred clinic location:***

**Beaverton**

OHSU Knight Cancer Institute  
15700 SW Greystone Court  
Beaverton, OR 97006

Phone number: 971-262-9000

Fax number: 503-346-8058

**NW Portland**

Legacy Good Samaritan campus  
Medical Office Building 3, Suite 150  
1130 NW 22nd Ave.  
Portland, OR 97210

Phone number: 971-262-9600

Fax number: 503-346-8058

**Gresham**

Legacy Mount Hood campus  
Medical Office Building 3, Suite 140  
24988 SE Stark  
Gresham, OR 97030

Phone number: 971-262-9500

Fax number: 503-346-8058

**Tualatin**

Legacy Meridian Park campus  
Medical Office Building 2, Suite 140  
19260 SW 65th Ave.  
Tualatin, OR 97062

Phone number: 971-262-9700

Fax number: 503-346-8058

Infusion orders located at: [www.ohsuknight.com/infusionorders](http://www.ohsuknight.com/infusionorders)