Denosumab (Prolia)

Provider Order Form



PATIENT INFORMATION

Dat	ate: Patient Name:			DOB:			
ICD-10 code (required):			ICD-10 description:				
□ NKDA Allergies:			Weight lbs/kg:				
Pat	ient Status: ☐ New to Therapy ☐ Continuing Ther	rapy N	Due Date ((if applicable):			
PR	OVIDER INFORMATION						
Ordering Provider:			Provider NPI:				
Referring Practice Name:			Phone:		Fax:		
Practice Address:			City:		State:	Zip Code:	
NU	JRSING	1	RAPY AD	OMINISTRATIO	ON		
0	including reaction management and post-procedure observation DEXA scan results and date (Please also attach results)		Denosumab (Prolia) Dose: □ 60mg/ml Route: subcutaneous injection Frequency: □ every 6 months				
Ø			Refills: □ Zero / □ One refill / □ Other: ———————————————————————————————————				

SPECIAL INSTRUCTIONS

Hypocalcemia: Must be corrected before initiating Prolia. May worsen, especially in patients with renal impairment. Adequately supplement patients with calcium and vitamin D.

Fax to 281-406-1047 or email: referrals@curbsideinfusion.com Office: 281-406-1046 Scheduling Ext 0 - scheduling@curbsideinfusion.com Billing Ext 1 - billing@curbsideinfusion.com www.curbsideinfusion.com

ADULT REACTION MANAGEMENT PROTOCOL

- Observe for hypersensitivity reaction: Fever, chills, rigors, pruritus, rash, cough, sneezing, throat irritation, nausea, vomiting.
- If reaction occurs:
 - If indicated, stop infusion.
 - Maintain/establish vascular access.
 - IVX Health clinicians have the following PRN medications available for the following reactions.
 - Headache, pain, fever > 100.4F, chills or rigors- Acetaminophen 650mg PO or Ibuprofen 400mg PO.
 - Rhinitis, allergies, hives, pruritis and other nonspecific symptoms of allergic reaction Loratadine 10mg PO or Diphenhydramine 25-50mg PO or IV
 - Nausea, vomiting, heartburn, acid reflux- Ondansetron 4mg ODT (may repeat x 1 in 20 minutes if nausea continues, max dose 8mg) or Famotidine 20mg PO.
 - Severe Nausea, vomiting, heartburn, acid reflux- Ondansetron 4mg SIVP (may repeat x 1 in 20 minutes if nausea continues, max dose 8mg) or Famotidine
 - Hypotension (90/60), vasovagal response- Place patient in reclined position, administer 0.9% Sodium Chloride IV 500ml. May repeat to keep BP >90/60, maximum of 1000ml, monitor vital signs.
 - Hypertension (>30 mmHg increase from baseline or >180 mmHg SBP): Clonidine 0.1mg and wait 45 minutes, may administer Amlodipine 5mg if hypertension persists
 - Chest pain/discomfort, shortness of breath- Oxygen 2-15 liters, titrate to keep Spo2 >92%.
 - Famotidine 20mg IV- Refractory to other treatments given
 - Solumedrol 125mg IV- Refractory to other treatments given.
 - When symptoms resolve resume infusion at 50% previous rate and increase per manufactures guidelines.
 - Notify referring provider as clinically appropriate and follow clinical escalation protocol.

Severe allergic/anaphylactic reaction:

- If symptoms are rapidly progressing or continuing after administration of prn medications above and signs symptoms of severe allergic/anaphylactic reaction (angioedema, swelling of the mouth, tongue, lips, or airway, dyspnea, bronchospasm with or without hypotension or hypertension).
 - Call 911
 - Initiate basic life support as needed. 0
 - Bring the **AED** to the patient (Attach pads if indicated). 0
 - Epinephrine- administer 0.3mg of a 1:1,000 (1mg/ml) concentration intramuscularly (preferably outer thigh), may be repeated every 5-15 minutes as needed to a maximum of 3 doses.
 - Place patient in recumbent position, elevate lower extremities.
 - Oxygen- administer 2-15 liters/minute or 100 percent oxygen as needed maintain SpO2 >92 percent.
 - IV Fluids- Treat hypotension with normal saline bolus of 500ml, repeat as needed to maintain systolic BP >90.
 - Administer diphenhydramine 50mg IV or Famotidine 20mg IVP, if not previously given.
 - Administer methylprednisolone 125mg IVP, if not previously given.
 - Continuous monitoring of blood pressure, pulse oximetry, and heart rate.
 - Notify clinical executive, DON or CMO, when appropriate. Must be done same day. Do not delay treatment.

Patient Name	Patient Date of Birth
Provider Name (Print)	
Provider Signature	Date

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page 2 of 2