

3/2023

Mepolizumab (NUCALA) Injection

NAME: **BIRTHDATE: INSURANCE: PROVIDER NAME: CLINIC NAME and Phone number:**

Patient identification

Moight	ka Hoighti em Allergiesi Diagnesis Codes
	kg Height:cm Allergies: Diagnosis Code:
	nt Start Date:
**If treat	tment not started within 90 days, new orders with updated date/signature will be required. Otherwise,
orders ar	re valid for 1 year from signature date.**
GUIDELIN	NES FOR ORDERING:
1)	Patient name and date of birth must be on EACH page of this order form. This form will be returned if any
	left blank.
2)	Send FACE SHEET and complete copy of INSURANCE information. Send H&P or most recent chart note,
	must be within the past 12 months.
3)	Send a copy of related lab results if not available for review in Epic.
MEDICA1	TIONS: (Must check one)
	Mepolizumab (NUCALA) injection, 40mg, subcutaneous, every 4 weeks.
	Mepolizumab (NUCALA) injection, 100mg, subcutaneous, every 4 weeks.
	Mepolizumab (NUCALA) injection, 300mg, subcutaneous, every 4 weeks.
	Mepolizumab (NUCALA) injection, mg, subcutaneous, every weeks.
HYPERSE	NSITIVITY INJECTION REACTION ORDERS
MILD REA	ACTION: Localized pruritus, rhinitis, localized rash, fever greater than 38 degrees C.
1. N	otify MD of symptoms and hypersensitivity reaction orders initiated.
MODERA	ATE REACTION: Generalized pruritus, generalized flushing, rash, back pain, mild dyspnea, chills,

- hypotension with SBP less than 90 mmHg or greater than 20% decrease in SBP.
 - 1. Start IV and maintain IV access with NS line per provider instruction. 2. Place patients in a supine position with lower extremities elevated unless respiratory distress or nausea/vomiting.
 - 3. Maintain airway.

SEVERE REACTION: Respiratory distress (bronchospasm, stridor, wheezing, persistent cough, respiratory depression, cardiac arrhythmia, chest pain, generalized urticaria (hives), syncope, gastrointestinal symptoms (abdominal pain, N/V, diarrhea), hypotension SBP less than 80 mmHg, face/throat/tongue swelling, shock, loss of consciousness.

- 1. Start IV and maintain IV access with NS line per provider instruction.
- 2. Stay with patient and have another nurse notify MD of symptoms and infusion reaction orders initiated.
- 3. Place patients in supine position with lower extremities elevated unless respiratory distress or nausea/vomiting.
- 4. Maintain airway.

MEDICATIONS:

- Acetaminophen (TYLENOL) tablet 650 mg, Oral, EVERY 4 HOURS PRN, Mild or moderate reaction. Starting when released, for 2 doses.
- DiphenhydraMINE (Benadryl) capsule, 25mg, PO, prn. Mild hypersensitivity reaction. May repeat x1.
- DiphenhydrAMINE (Benadryl) injection, 25 mg, Intravenous, EVERY 15 MIN PRN, Moderate or severe reaction, may repeat x1. Starting when released, For 2 doses.
- PrednisONE (Deltasone) tablet, 40mg, PO. Once, for Mild hypersensitivity reaction.



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- Methylprednisolone sodium succinate (solu-MEDROL) 62.5mg/ml injection, 125 mg, Intravenous, ONCE PRN, moderate or severe reaction. Starting when released, For 1 dose. Mix with 2 mL provided diluent to make 62.5 mg/mL.
- Famotidine (Pepcid) tablet, 20mg, PO. Once, for Mild hypersensitivity reaction.
- Famotidine (Pepcid) injection 20 mg, Intravenous, ONCE PRN, Moderate or severe reaction, for 1 dose. Dilute 2 mL of famotidine with 8 mL of normal saline to a final concentration of 2 mg/mL. Administer ordered dose over a period of at least 2 minutes.
- Albuterol 90 mcg/puff inhaler 2 puff, Inhalation, ONCE PRN, Shortness of breath, severe reaction. Starting when released, For 1 dose. Shake well. Use with spacer.
- Start oxygen. For *MODERATE* Reaction: Start 2 liters O2 per nasal cannula as needed for SpO2 less than 93%. For *SEVERE* Reaction: Start continuous pulse oximetry. Administer 6 to 8 liters per minute (LPM) via face mask, or up to 100% oxygen to maintain oxygen saturation greater than 90%.
- EPINEPHrine 1 mg/mL injection 0.3 mg. Intramuscular, EVERY 5 MIN PRN, Anaphylaxis. Starting when released, For 3 doses. Administer in the anterior lateral thigh using 1 1.5-inch needle for airway obstruction symptoms and/or hypotension.
- Sodium chloride 0.9% (NS) infusion 500ml. At 500 mL/hr., Intravenous, PRN, Moderate reaction, or anaphylaxis. Starting when released. Run wide open to gravity.

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.

My physician license Number is #	and I am acting within my scope of practice and
authorized by law to order infusion of the med identified on this form.	lications and blood products described above for the patient
Provider's printed name:	
Provider's signature:	
Date:	



ADULT AMBULATORY INFUSION ORDER 3/2023

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Please o	complete the following intake qu	estions.
Mobilit	y status	
	Independent	
	Walker	
	Wheelchair	
	Fully dependent	
Bariatri	c needs	
	Yes, briefly describe	
	No	
Transpo	ortation concerns	
	Yes, briefly describe	
	No	
Interpre	eter Service required	
	Yes, language	
	No	
	ient Infusion Services Intake Tear	
Please o	check the appropriate box for the	e patient's preferred infusion center location:
	□ PORTLAND	Phone- 503-215-6046 Fax- 503-487-3582
	□ WILLAMETTE FALLS	
	☐ MEDFORD	Phone- 541-732-7000 Fax- 541-732-3939
	☐ HOOD RIVER	Phone- 541-387-6445 Call for fax #
	□ SEASIDE	Phone- 503-717-7650 Fax- 503-470-3146
	☐ NEWBERG	Phone- 503-537-6040 Fax- 503-205-8720