



PATIENT INFORMATION (Complete or Fax Existing Chart)	PRESCRIBER INFORMATION
Name: _____ DOB: _____	Prescriber Name: _____
Address: _____	State License: _____
City, State, Zip: _____	NPI #: _____ DEA: _____
Phone: _____ Alt. Phone: _____	Address: _____
Email: _____ SS#: _____	City, State, Zip: _____
Gender: <input type="checkbox"/> M <input type="checkbox"/> F Weight: _____ (lbs) Ht: _____	Phone: _____ Fax: _____
Allergies: _____	Office Contact: _____ Phone: _____

INSURANCE INFORMATION – AND – Send a copy of the patient's prescription/insurance cards (front & back)

Primary Insurance: _____	Secondary Insurance (If Applicable): _____
Plan #: _____	Plan #: _____
Group #: _____	Group #: _____
RX Card (PBM): _____	RX Card (PBM): _____
BIN: _____ PCN: _____	BIN: _____ PCN: _____

CLINICAL INFORMATION

J45.50 Severe persistent asthma, uncomplicated J45.51 Severe persistent asthma with (acute) exacerbation J82.83 Eosinophilic asthma

J33.0 Polyp of the nasal cavity M30.1 Polyarteritis with lung involvement [Churg-Strauss] Other: _____

Prior Anaphylactic Reaction: No Yes (Reason/Date): _____

Other Medications: _____

Lab Results:

Positive Skin or RAST test to Perennial Aeroallergen: Yes No Test Date: _____ - _____ - _____

Serum IgE Level _____ IU/ML Test Date: _____ - _____ - _____

Serum Eosinophil Level: _____ cells/mcL Test Date: _____ - _____ - _____

Sputum Eosinophiles _____ cells/mcL Test Date: _____ - _____ - _____

NUCALA® ORDERS

Prescription type: New start Restart Continued therapy Total Doses Received: _____ Date of Last Injection: _____

Medication	Directions	Quantity/Refills
<input type="checkbox"/> Nucala (mepolizumab) 100mg/mL	<input type="checkbox"/> Inject 100mg under the skin once every 4 weeks. <input type="checkbox"/> Inject 300mg (3 separate 100mg injections) under the skin once every 4 weeks. <input type="checkbox"/> Other: _____	Quantity: _____ Refills: _____
Pre-Medication	Dose/Strength	Directions
<input type="checkbox"/> Acetaminophen	<input type="checkbox"/> 500mg	<input type="checkbox"/> Take 1-2 tablets PO prior to infusion or post-infusion as directed
<input type="checkbox"/> Diphenhydramine	<input type="checkbox"/> 25mg IV/PO <input type="checkbox"/> 50mg IV/PO	<input type="checkbox"/> Take 1 tablet PO prior to infusion or as directed OR <input type="checkbox"/> Inject contents of 1 vial IV prior to infusion or as directed
<input type="checkbox"/> Methylprednisolone	<input type="checkbox"/> 40mg <input type="checkbox"/> 100mg <input type="checkbox"/> 125mg	<input type="checkbox"/> Inject contents of 1 vial IV prior to infusion or as directed <input type="checkbox"/> Other: Inject 100mg IV 30 minutes prior to infusion
<input type="checkbox"/> _____	_____	_____

INFUSION REACTION ORDERS

CONFIDENTIALITY STATEMENT: This facsimile and documents accompanying this transmission contain confidential health information that is legally privileged. This information is intended only for the use of the individual or entity named above. The authorized recipient of this information is prohibited from disclosing this information to any other party unless required to do so by law or regulation. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution, or action taken in reliance on the contents of these documents is strictly prohibited. If you have received this information in error, please notify the sender at the address and telephone number set forth herein and arrange for return or destruction of the material. In no event should such material be read by anyone other than the named addressee, except by express authority of the sender to the named addressee.



Mild reaction protocol:

- Diphenhydramine 25mg IV, one time, for pruritus.
- If symptoms worsen, see orders for moderate to severe reactions.*

Moderate reaction protocol:

- Acetaminophen 650mg PO, one time, for pyrexia or rigors
- Diphenhydramine 50mg IV, one time, for pruritus or urticaria
- Methylprednisolone 125mg IV, one time, for respiratory or neurologic symptoms
- If symptoms worsen, see interventions for severe reactions*

Severe reaction protocol: (Call 911 if initiated):

- Titrate oxygen via continuous flow per nasal cannula or face mask to maintain spO2 of greater than ninety-five percent (>95%)
- Diphenhydramine 50mg IV, one time, for respiratory symptoms, edema, or anaphylaxis
- Methylprednisolone 125mg IV, one time, for respiratory symptoms, edema, or anaphylaxis
- Sodium Chloride 0.9% 500mL IV over 30-60 min, one time, for cardiovascular symptoms
- Epinephrine 0.3mg/0.3mL IM into mid-antrolateral aspect of thigh of anaphylaxis, may repeat x1 in 5-15 minutes if symptoms are not resolved or worsen

FLUSHING & LOCKING ORDERS

Flushing Protocol (>66lbs/33kg)

PIV and Midline:

- 0.9% Sodium Chloride 2-5mL IV flush before and after each infusion

Implanted Port, PICC, Tunneled Catheter, and Non-tunneled Catheter:

- 0.9% Sodium Chloride 5mL IV flush before infusion/lab draw and 10mL IV flush after infusion/lab draw

Locking Protocol (>66lbs/33kg)

PIV and Midline:

- Heparin Sodium 10 units/mL 1mL IV final flush post normal saline flush

PICC:

- Heparin Sodium 10 units/mL 3mL IV final flush post normal saline flush

Implanted Port, Tunneled Catheter, and Non-tunneled Catheter:

- Heparin Sodium 100 units/mL 3-5mL IV final flush post normal saline flush

**** May substitute Dextrose 5% in Water, or alternative, for 0.9% Sodium Chloride, when indicated due to incompatibility with medications being infused**

SIGNATURE

We hereby authorize Infuse IQ LLC to provide all supplies and additional services (nursing/patient training) required to provide and deliver the medicine as prescribed in this referral.

X _____

Prescriber Signature

Date: _____

To ensure payment by insurance carrier, please include supporting clinical documentation for specified ICD 10 Code, demographic, and insurance information along with faxed order. Initial appointment will be verified upon insurance approval.

CONFIDENTIALITY STATEMENT: This facsimile and documents accompanying this transmission contain confidential health information that is legally privileged. This information is intended only for the use of the individual or entity named above. The authorized recipient of this information is prohibited from disclosing this information to any other party unless required to do so by law or regulation. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution, or action taken in reliance on the contents of these documents is strictly prohibited. If you have received this information in error, please notify the sender at the address and telephone number set forth herein and arrange for return or destruction of the material. In no event should such material be read by anyone other than the named addressee, except by express authority of the sender to the named addressee.