

Referral Status:		MRN:	
<input type="checkbox"/> New referral	<input type="checkbox"/> Order change	<input type="checkbox"/> Order Renewal	
Patient preferred clinic:			

Xolair[®] (omalizumab) Standard Plan of Treatment for Nasal Polyps
PATIENT DEMOGRAPHICS:

Date of Referral:	Patient's Phone:				
Patient Name:	Address:				
Date of Birth:	City, State, Zip:				
Height in inches:	Weight: LB or KG	Gender:	Allergies:	<input type="checkbox"/> See list	<input type="checkbox"/> NKDA

DIAGNOSIS: (PLEASE COMPLETE 2ND AND 3RD DIGITS TO COMPLETE ICD 10 FOR BILLING)

<input type="checkbox"/> J33.0 - Polyp of nasal cavity	<input type="checkbox"/> J33.1 - Polypoid sinus degeneration
<input type="checkbox"/> J33.8 - Other polyp of sinus	<input type="checkbox"/> J33.9 - Nasal polyp, unspecified
<input type="checkbox"/> - Other:	

REQUESTED DOCUMENTATION:

1 Insurance information	IF NO:	IF YES:
2 Most recent History & Physical	PLEASE STATE	LAST INFUSION DATE:
3 Full medication list	REQUIRED WASHOUT	NEXT INFUSION DATE:
4 Tried and failed therapies	FROM PREVIOUS	IF ORDER CHANGE:
5 Pre-treatment serum IgE level as required for pretreatment dosing	THERAPY:	
		<input type="checkbox"/> Continue current order until insurance approved

Provider Attestation for HCP administration:

<input type="checkbox"/> Provider attests that the patient or caregiver is not competent or is physically unable to administer the Xolair labeled self-administration.	<input type="checkbox"/> The location and circumstances for self-administration are not adequate for the potential treatment of anaphylaxis should that arise.
<input type="checkbox"/> Patient has experienced severe hypersensitivity reactions to Xolair or other agents, such as foods, drugs, biologics, within the past 6 months or requires administration and direct monitoring by a healthcare professional.	<input type="checkbox"/> Patient has a history of uncontrolled disease and ordering physician attests that in their clinical opinion, it is not advisable to try the self-administration formulation of requested drug
<input type="checkbox"/> Patient has not received at least 3 doses of Xolair under the guidance of a healthcare provider with no hypersensitivity reactions*	<input type="checkbox"/> Due to patient's weight, ordering provider attests that in their clinical opinion, it is not advisable to try the self-administered formulation of requested drug.

*Specific reactions: _____

MEDICATION ORDERS:
NOTE: Patient may be ineligible to receive Xolair[®] (omalizumab) if patient has signs/symptoms of parasitic infection, is currently being treated for a parasitic infection, or is having acute bronchospasm and/or asthma attack.
MEDICATION/FREQUENCY:
 Xolair[®] (omalizumab) subcutaneously every 2 weeks:
 Xolair[®] (omalizumab) subcutaneously every 4 weeks:

DOSE:
 75mg/dose
 150 mg/dose
 225mg/dose
 300mg/dose
 375mg/dose
 450mg/dose
 525mg/dose
 600mg/dose

Administer as subcutaneous injection to upper arm, thigh, or abdomen.

SPECIAL ORDERS:

POST WAIT: *Extended post treatment monitoring for any patient new to therapy*

Standard Palmetto Infusion Post wait per package insert: Monitor patient for two (2) hours after first injection, for (1) hour after second injection, for 30 minutes after third injection, then monitor for 15-minutes with all subsequent injections. Unless otherwise selected below.

 Monitor patient for two (2) hours after first 3 injections, and for 30-minutes after all subsequent injections.
 Provider specific post wait: _____

 Refills x 12 months unless noted otherwise here:

CARE ORDERS:
 Provide nursing care per Palmetto Infusion Nursing Procedures and post procedure observation if indicated

ADVERSE REACTION & ANAPHYLAXIS ORDERS:

Administer acute infusion and anaphylaxis medications per Palmetto Infusion standing adverse reaction orders, which can be found at our website or scan here.


PRESCRIBER INFORMATION:

PROVIDER NAME:	PHONE:
ADDRESS:	FAX:
CITY, STATE, ZIP:	NPI:

PRESCRIBER SIGNATURE: (No stamp signatures)
DATE

_____	_____
Dispense as written/Brand medically necessary	Substitution permitted