

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 Urticaria

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"/> L50.0 - Allergic Urticaria	<input type="checkbox"/> L50.1 - Chronic Idiopathic Urticaria (CIU)
<input type="checkbox"/> L50.8 - Other Specified Urticaria	<input type="checkbox"/> - Other: _____

REQUESTED DOCUMENTATION:

1	Insurance information
2	Most recent History & Physical
3	Full medication list
4	Tried and failed therapies
5	Pre-treatment serum IgE level as required for pre treatment dosing

PREVIOUS ADMINISTRATION: HAS THIS PATIENT TAKEN THIS MEDICATION BEFORE?

IF NO:	IF YES:
PLEASE STATE REQUIRED WASHOUT FROM PREVIOUS THERAPY:	LAST INFUSION DATE:
	NEXT INFUSION DATE:
IF ORDER CHANGE:	
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 4 weeks: Other: _____

DOSE:

150 mg/dose 300mg/dose

Administer as subcutaneous injection to upper arm, thigh, or abdomen. No more than 150 mg per injection site.

SPECIAL ORDERS:

Extended post treatment monitoring for any patient new to therapy: monitor patient for two (2) hours after first 3 injections, and for 30-minutes after all subsequent injections.

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