

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

Xolair® (omalizumab) Standard Plan of Treatment for IgE-Mediated Food Allergy

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"/> NDKA

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

<input type="checkbox"/> Z91.010 Allergy to peanuts	<input type="checkbox"/> Z91.013 Allergy to seafood
<input type="checkbox"/> Z91.011 Allergy to milk products	<input type="checkbox"/> Z91.018 Allergy to other foods
<input type="checkbox"/> Z91.012 Allergy to eggs	
<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 dosing	THERAPY:	
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
 400mg/dose 450mg/dose 525 mg/dose 600mg/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 injection, for one (1) hour after second injection, for 30-minutes after third injection, and then 15-minutes after all subsequent injections.

Refills x 12 months unless noted otherwise here:

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