

Dispense as written/Brand medically necessary

Referral Status:	MRN:	
New referral	Order change	Order Renewal
Patient preferred clinic:		

Substitution permitted

Phone: 1-800-809-1265 Fax: 1-866-872-8920 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: IB or KG Gender: Allergies: Height in inches: Weight: NDKA See list DIAGNOSIS: (PLEASE COMPLETE 2ND AND 3RD DIGITS TO COMPLETE ICD 10 FOR BILLING) J33.0 Polyp of nasal cavity J33.1 Polypoid sinus degeneration J33.9 Nasal polyp J33.8 Other polyp of sinus - Other: **REQUESTED DOCUMENTATION:** PREVIOUS ADMINISTRATION: HAS THIS PATIENT TAKEN THIS MEDICATION BEFORE? Insurance information IF NO: IF YES: PLEASE STATE LAST INFUSION DATE: Most recent History & Physical REQUIRED WASHOUT Full medication list NEXT INFUSION DATE: FROM PREVIOUS Tried and failed therapies IF ORDER CHANGE: THERAPY: Pre-treatment serum IgE level as required for Continue current order until insurance approved pretratment dosing **Provider Attestation for HCP administration:** Provider attests that the patient or caregiver is not competent or is physically The location and circumstances for self-administration are not adequate for the unable to administer the Xolair labeled self-administration potential treatment of anaphylaxis should that arise. Patient has experienced severe hypersensitivity reactions to Xolair or Patient has a history of uncontrolled disease and ordering physician attests that in their other agents, such as foods, drugs, biologics, within the past 6 months or requires clinical opinion, it is not advisable to try the self-administration formulation of requested administration and direct monitoring by a healthcare professional. Patient has not received at least 3 doses of Xolair under the guidance of a Due to patient's weight, ordering provider attests that in their clinical opinion, it is not healthcare provider with no hypersensitivity reactions* 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. 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 30minutes after third injection, and then 15-minutes for 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