

PATIENT INFORMATION
PATIENT NAME: _____ **PHONE:** _____

DATE OF BIRTH: _____ **SEX:** ☐ M ☐ F **HEIGHT:** _____ **WEIGHT:** _____ ☐ LBS ☐ KG

ALLERGIES: _____ **PREFERRED CLINIC:** _____

REFERRAL STATUS: ☐ NEW REFERRAL ☐ ORDER CHANGE ☐ ORDER RENEWAL

DIAGNOSIS & CLINICAL DOCUMENTATION
***PLEASE COMPLETE ICD-10 FOR SPECIFIC DIAGNOSIS**
☐ J33. ___ Nasal Polyps

☐ J45. ___ Persistent Asthma

☐ L50. ___ Urticaria

☐ Z91. ___ Food Allergy

☐ ICD-10 CODE: _____ DESCRIPTION: _____

REQUIRED DOCUMENTATION
☐ Insurance Information

☐ List of Medications

☐ Tried & Failed Therapies

☐ Most Recent History & Physical

☐ Pre-Treatment IgE levels

☐ Skin or RAST Test, if applicable

MEDICATION ORDER
***Please complete dose when selecting frequency**
☐ Xolair® (omalizumab) _____ mg
via subcutaneous injection every 2 weeks

☐ Xolair® (omalizumab) _____ mg
via subcutaneous injection every 4 weeks

☒ REFILL X 12 MONTHS UNLESS OTHERWISE NOTED
HERE: _____

Provider Attestation of need for HCP administration

INITIALS	Patient has experienced severe hypersensitivity reactions to Xolair within the past 6 months and requires administration and direct monitoring by a healthcare professional.
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INITIALS	Patient or caregiver are not competent or are physically unable to administer the Xolair formulation labeled for self-administration.
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INITIALS	Patient has history of uncontrolled disease and in the clinical opinion of the ordering provider, it is not advisable to try the self-administered formulation of Xolair.
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INITIALS	The circumstances and location for self-administration are not adequate for the potential treatment of anaphylaxis.
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INITIALS	Patient's weight is such that in the clinical opinion of the ordering provider, it is not advisable to try the self-administered formulation of Xolair.
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Patient to be observed for 30 minutes following the first injection and then for 15 minutes following each subsequent injection.
In the event of an adverse reaction occurring in the infusion clinic, utilize the Immersiv Health adverse reaction protocol.

PRESCRIBER INFORMATION
PROVIDER NAME: _____ **NPI #:** _____

EMAIL: _____ **PHONE:** _____ **FAX:** _____

ADDRESS (INCLUDE CITY, STATE, ZIP): _____

SUPERVISING PHYSICIAN: _____ **CONTACT NAME:** _____
(IF APPLICABLE)
SIGNATURE: _____ **DATE:** _____
(NO STAMPS)

SUBSTITUTION PERMITTED

DISPENSE AS WRITTEN