

Patient Name: \_\_\_\_\_ DOB: \_\_\_\_\_ Date of Last Infusion: \_\_\_\_\_  
Height \_\_\_\_\_ Weight \_\_\_\_\_  
Infusion Location: (state and site) \_\_\_\_\_

## Xolair<sup>®</sup> (omalizumab) Orders

Diagnosis (please provide ICD-10 code in space provided):

\_\_\_\_\_ Allergic asthma  
(ICD-10)

\_\_\_\_\_ Chronic spontaneous urticaria  
(ICD-10)

\_\_\_\_\_ Other: \_\_\_\_\_  
(ICD-10)

Provider: Check to confirm patient has received epinephrine auto-injector & has been educated on use

IgE Result/Date: \_\_\_\_\_

### Nursing Orders:

-Hold treatment and notify provider if patient:

- Reports signs or symptoms of serum sickness (fever, rash, joint pain/swelling/stiffness, muscle pain, swollen lymph nodes)
- Has not received epinephrine auto-injector and education on its use

Xolair \_\_\_\_\_ mg subcutaneously. Divide doses exceeding 150 mg among multiple injection sites to limit injections to not more than 150 mg per site

Dosing:

For Asthma/Chronic Rhinosinusitis dose based on IgE levels and weight

For Chronic Spontaneous Urticaria flat dose of 150mg or 300mg

Frequency: Every \_\_\_\_\_ weeks

### Observation Period:

- Following the first three injections, monitor patient for post-injection observation period of 2 hours. For all subsequent injections, monitor patient for 30 minutes
- Record vital signs prior to discharge
- If patient develops bronchospasm, angioedema, hypotension, urticaria or other signs of anaphylaxis, initiate Hypersensitivity Reaction Management Protocol to include administration of epinephrine 0.3 mg IM STAT

### Additional Orders:

Provider Name (print): \_\_\_\_\_ Date: \_\_\_\_\_

Provider Signature: \_\_\_\_\_ Time: \_\_\_\_\_