

### Standard Plan of Treatment for NULOJIX

(Re)Certification Period from \_\_\_\_\_ to \_\_\_\_\_

**NOTE:** Patient is *ineligible* to receive Benlysta if receiving antibiotic for active infectious process (due to the possibility of developing a super-infection related to its effect on the immune status), or if he/she has a suspected infection.

Patient's Name: \_\_\_\_\_ Weight: \_\_\_\_\_

Allergies: \_\_\_\_\_ Height: \_\_\_\_\_

Primary Diagnosis \_\_\_\_\_ Secondary Diagnosis: \_\_\_\_\_

**NULOJIX is contraindicated in transplant recipients who are EBV seronegative or unknown serostatus.**

#### Orders:

Obtain weight each visit. Vital Signs: baseline and every 30 minutes until infusion complete.  
Instruct patient/caregiver on medications, signs/symptoms of adverse reaction. Assess patient response to therapy.  
Utilize existing central line for administration, or initiate a peripheral IV with each infusion, prn.  
Normal Saline Flush 3-10 ml before infusion, after primary drug has infused, Infuse Normal Saline 0.9% or D5W 20-50 ml to flush tubing/line, followed by Heparin Lock 1-5ml 100 units/ml as needed per line type.  
Pump, tubing, and supplies needed to complete prescribed therapy.  
**If adverse drug reaction, implement the Standing Adverse Reaction protocol.**

#### Drug:

Weight on Transplant \_\_\_\_\_ (*Dose is calculated on transplant weight unless weight varies by  $\geq 10\%$* )

**Nulogix 5mg/kg IV in 100ml NS, administer over 30 minutes every 4 weeks**

Labs: \_\_\_\_\_

Pharmacist to perform clinical drug monitoring.

(No Stamped Signatures please)

Physician's Signature: \_\_\_\_\_ / \_\_\_\_\_ Date: \_\_\_\_\_

(Dispense as written)

(Substitution permitted)

Print Physician Name: \_\_\_\_\_

**PLEASE FAX DEMOGRAPHICS AND INSURANCE INFORMATION TO 866-872-8920**