

Referral Status:  New Patient  Updated Patient Information

## CMC Infusion Center- Referral Face Sheet



Patient Name: \_\_\_\_\_ Patient D.O.B. \_\_\_\_\_

Current Height: \_\_\_\_\_ Current Weight \_\_\_\_\_

Patient Phone # (\_\_\_\_\_) \_\_\_\_\_ - \_\_\_\_\_ Gender \_\_\_\_\_

Patient Address \_\_\_\_\_

City \_\_\_\_\_ State \_\_\_\_\_ Zip Code \_\_\_\_\_

Related Diagnosis Code (ICD-10 code): \_\_\_\_\_

Referring Provider (Please Print) \_\_\_\_\_

Office Phone # (\_\_\_\_\_) \_\_\_\_\_ - \_\_\_\_\_ Office Contact: \_\_\_\_\_

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**If referring provider and/or patient is external to Conway Medical Center. Please include:**

- Most recent history and physical including:
  - Comprehensive medication list
  - Past medical history
  - Related past/ failed therapies (with dates)
  - Known allergies
- Most recent labs / test result (Ex. CBC, PPD, Hepatitis screening, etc.)
- Insurance Information

Please fax requested documentation, face sheet, and treatment plan to **843-234-5460** attention CMC Medication Management Specialist. Please feel free to call 843-234-8575 with any questions.

We look forward to the opportunity to be a part of your patient's treatment journey at Conway Medical Center.

Referral Status  New  Order Change  Order Renewal



# Actemra® (tocilizumab) Treatment Plan

Patient Name: \_\_\_\_\_ Patient DOB \_\_\_\_\_

### General

- Diagnosis- Rheumatoid Arthritis Seropositive multiple sites ICD-10 Code \_\_\_\_\_
- Diagnosis- Rheumatoid Arthritis Seronegative multiple sites ICD-10 Code \_\_\_\_\_
- Diagnosis- Giant cell arteritis ICD-10 Code \_\_\_\_\_

Pt. Weight \_\_\_\_\_ Pt Height \_\_\_\_\_ Known Allergies \_\_\_\_\_

Requested Start Date \_\_\_\_/\_\_\_\_/\_\_\_\_ Requested Frequency \_\_\_\_\_ weeks

Has patient previously received this medication  No  Yes, if so, date of last infusion \_\_\_\_/\_\_\_\_/\_\_\_\_

Authorized Treatment Duration  12 Months  Other \_\_\_\_\_

Infusion (Month 1 to 12) Duration: 12 Months

- Vital Signs (Month 1 to 12)  
*every 30 min, Resp, BP, HR, Temperature on admission to the unit and prior to infusion HR, BP every 30 minutes during the infusion and prior to discharge*

Tylenol (Month 1 to 12)

- 650 mg, Oral, Tab, Day of Tx  
**Comments: Administer 30 minutes before tocilizumab dose**

Benadryl (Month 1 to 12)

- 25 mg, IV Push, Injection, Day of Tx  
**Comments: Administer 30 minutes before tocilizumab dose**

- 50 mg, IV Push, Injection, Day of Tx  
**Comments: Administer 30 minutes before tocilizumab dose**

SOLU-Medrol (Month 1 to 12)

- 40 mg, IV Push, Powder-Inj, Day of Tx  
**Comments: Administer 30 minutes before tocilizumab dose**

Actemra (Month 1 to 12)

- 4 mg/kg, IV Piggyback, Injection  
*Comments: infuse in 100 mL 0.9% NaCl over 60 minutes. Nurse to slowly flush tubing with up to 20 mL 0.9% NS once infusion is finished to clear all medication in IV tubing. Max dose = 800 mg*

- 8 mg/kg, IV Piggyback, Injection,  
*Comments: infuse in 100 mL 0.9% NaCl over 60 minutes. Nurse to slowly flush tubing with up to 20 mL 0.9% NS once infusion is finished to clear all medication in IV tubing. Max dose = 800 mg*

- Communication Order (Month 1 to 12)  
*Patient must stay in infusion area for 30 min post infusion. If vital signs normal may be discharged to home. If abnormal, call physician. Patient should be accompanied by another adult upon discharge*

### Prescriber Signature (No Stamped Signatures or Electronic Signatures)

Provider Signature \_\_\_\_\_ Provider Name (please print) \_\_\_\_\_

Date \_\_\_\_/\_\_\_\_/\_\_\_\_ Patient Name \_\_\_\_\_ Patient DOB \_\_\_\_\_

# Actemra® (tocilizumab) Treatment Plan

- Communication Order (Month 1 to 12)  
*Ensure the CBC with diff is available within 2 months of initiation or within 3 months of subsequent infusions of tocilizumab (Actemra). Notify provider if labs are not within defined parameters: Note Defined Parameters=: absolute neutrophil count (ANC) of 2000/mm<sup>3</sup> or greater; platelet count of 100,000/mm<sup>3</sup> or greater are required before initiating tocilizumab (Actemra)*
- Communication Order (Month 1 to 12)  
*Ensure AST/ALT results are available within 2 months of initiation or within 3 months of subsequent infusions. Notify provider if results outside of normal limits: Note: Do **not** initiate tocilizumab in patients with baseline ALT or AST levels greater than 1.5 x ULN*
- Communication Order (Month 1 to 12)  
*Ensure negative PPD or other test to exclude latent tuberculosis. Results must be within the last 12 months.*
- Notify Provider (Month 1 to 12)  
*for fever/chills, chest pain, hypotension, hypertension, dyspnea, Pruritis, urticaria, persistent flushing, Temperature > 100, or HR <50 or > 130*
- Notify Provider (Month 1 to 12)  
*For POSITIVE results of PPD test or other test to exclude latent tuberculosis. Results must be within the last 12 months.*
- INF Infusion Room Orders Subphase (Month 1 to 12)
- heparin flush 100 units/mL intravenous solution (Month 1 to 12)  
*5 mL, Intracatheter, Injection, Once, PRN other (see comment)  
Comments: To maintain central line patency for implanted ports, flush once monthly or prior to deaccess*

## Labs (Month 1 to 12) Duration: 12 Months

- QuantiFERON TB Gold Plus (if needed)
- CBC w/ Diff (Month 1 to 12)  
*Blood, Routine, (if needed)*
- AST (Month 1 to 12)  
*Blood, Routine, (if needed)*
- ALT (Month 1 to 12)  
*Blood, Routine, (if needed)*
- Sedimentation Rate (Month 1, 3, 6, 9, 12)  
*Blood, Routine, (if needed)*
- BUN (Month 1, 3, 6, 9, 12)  
*Blood, Routine, (if needed)*
- Creatinine w/GFR (Month 1, 3, 6, 9, 12)  
*Blood, Routine, (if needed)*
- Lipid Panel (Month 1, 6, 12)  
*Blood, Routine, (if needed)*

## Prescriber Signature (No Stamped Signatures or Electronic Signatures)

Provider Signature \_\_\_\_\_ Provider Name (please print) \_\_\_\_\_

Date \_\_\_\_/\_\_\_\_/\_\_\_\_ Patient Name \_\_\_\_\_ Patient DOB \_\_\_\_\_

Biosimilar Substitution **NOT** permitted