

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



# Ocrevus (ocrelizumab) Treatment Plan

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

### General

Diagnosis- \_\_\_\_\_ ICD-10 Code \_\_\_\_\_

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

Requested Start Date \_\_\_\_/\_\_\_\_/\_\_\_\_

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*

**Pre- Treatment Medications** (Note: Pre-treatment with acetaminophen oral, diphenhydramine IV push, and methylprednisolone IV push recommended prior to infusion)

Acetaminophen (Month 1 to 12)

1000 mg, Oral, Tab, Day of Tx  
**Comments: Administer 30 minutes before dose**

Diphenhydramine (Month 1 to 12)

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

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

Methylprednisolone (Month 1 to 12)

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

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

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

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

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

# Ocrevus (ocrelizumab) Treatment Plan

## Loading Dose Ocrelizumab

- 300 mg, IV Piggyback, Injection, **Once**  
*Comments: 300 mg dose: Begin infusion at 30 mL/hour; increase by 30 mL/hour every 30 minutes to a maximum rate of 180 mL/hour. Infusion duration is 2.5 hours or longer. Monitor for infusion reactions during infusion and observe for at least one hour after infusion is complete. If infusion reaction occurs, interrupt infusion, discontinue or decrease the rate, depending on the severity of the reaction and contact provider*
- 300 mg, IV Piggyback, Injection, 2 doses on **Day 1 and day 14**  
*Comments: 300 mg dose: Begin infusion at 30 mL/hour; increase by 30 mL/hour every 30 minutes to a maximum rate of 180 mL/hour. Infusion duration is 2.5 hours or longer. Monitor for infusion reactions during infusion and observe for at least one hour after infusion is complete. If infusion reaction occurs, interrupt infusion, discontinue or decrease the rate, depending on the severity of the reaction and contact provider*

## Maintenance Dose Ocrelizumab

- 600 mg, IV Piggyback, Injection, **Once**  
*Comments: 600 mg dose: Begin infusion at 40 mL/hour; increase by 40 mL/hour every 30 minutes to a maximum rate of 200 mL/hour. Infusion duration is 3.5 hours or longer. Monitor for infusion reactions during infusion and observe for at least one hour after infusion is complete. If infusion reaction occurs, interrupt infusion, discontinue or decrease the rate, depending on the severity of the reaction and contact provider*
- 600 mg, IV Piggyback, Injection, **Every 6 months starting** \_\_\_/\_\_\_/\_\_\_  
*Comments: 600 mg dose: Begin infusion at 40 mL/hour; increase by 40 mL/hour every 30 minutes to a maximum rate of 200 mL/hour. Infusion duration is 3.5 hours or longer. Monitor for infusion reactions during infusion and observe for at least one hour after infusion is complete. If infusion reaction occurs, interrupt infusion, discontinue or decrease the rate, depending on the severity of the reaction and contact provider*

- Communication Order  
*Administer through a dedicated IV line using a 0.2 or 0.22 micron in-line filter*
- Communication Order  
*Constant order, If patient develops fever, chills, rigors, hypotension or respiratory difficulty during infusion, STOP infusion and call MD/LIP.*
- Communication Order (Month 1 to 12)  
*Patient must stay in infusion area for 60 min post infusion. If vital signs normal may be discharged to home. If abnormal, call physician.*
- Communication Order (Month 1 to 12)  
*Ensure results of Hepatitis B surface antigen testing and record results if not present in medical record. Results must be within last 12 months.*
- 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*

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

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

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

# Ocrevus (ocrelizumab) Treatment Plan

*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.*
- Notify Provider (Month 1 to 12)  
*For POSITIVE results of Hepatitis B surface antigen or if results are not available. Results must be within the last 12 months.*
- INF Infusion Room Orders Subphase (Month 1 to 12)

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

- Quantiferon(R)-TB Gold, 4T, Incu-(36971) (If Needed)
- Hepatitis B Surface Antibody, Qual-(499) (If Needed)
- Hepatitis B Surface Antigen w Ref/Conf-(498) (If Needed)
- Hepatitis B Core Antibody, Total-(501) (If Needed)
- CBC w/ Diff (If Needed)
- Sedimentation Rate (If Needed)
- C-Reactive Protein (If Needed)
- BUN (If Needed)
- ALT(If Needed))
- AST (If Needed)
- BMP (If Needed)
- Urinalysis with Microscopic, if indicated (If Needed)

### Patient Monitoring Acknowledgement:

\* CMC Outpatient Infusion Services (OIS) shall ensure TB testing and Hepatitis B testing is completed prior to therapy initiation. If positive, OIS team will notify referring provider. Referring provider to manage subsequent care coordination for patients who screen TB positive, HBsAg+, or HBcAB+, or have increasing viral load.

Provider Initials \_\_\_\_\_

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

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

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