

Phone: 1-800-809-1265 Fax: 1-866-872-8920

MRN: \_\_\_\_\_

DOB: \_\_\_\_\_

## STANDARD Rituxan® (rituximab) PLAN OF TREATMENT

**NOTE:** Patient may be ineligible to receive rituximab if receiving antibiotics for active infectious process, antifungal therapy, active fever and/or suspected infection, newly diagnosed cardiac arrhythmias, severe abdominal pain or vomiting, and/or surgery.

**1. Patient Name:** \_\_\_\_\_ Height (inches): \_\_\_\_\_ Weight (lbs): \_\_\_\_\_

**2. Allergies:** \_\_\_\_\_

**3. Diagnosis:** \* Please complete the 2<sup>nd</sup> and 3<sup>rd</sup> digits to complete the ICD-10 code for billing

**M05.** \_\_\_\_\_ Rheumatoid Arthritis with Rheumatoid factor  **M06.** \_\_\_\_\_ Rheumatoid Arthritis without Rheumatoid factor

Other **ICD-10 Code:** \_\_\_\_\_ Diagnosis description: \_\_\_\_\_

**4. Pre-medications:** Administered 30 minutes prior to infusion as selected:

\* Premedication of Acetaminophen PO, Diphenhydramine IVP, and Methylprednisolone IVP is suggested prior to infusion

**Acetaminophen:**

- 650 mg PO
- 500 mg PO
- 325 mg PO
- 1000 mg PO

**Diphenhydramine:**  25 mg IVP,  50 mg IVP,  25 mg PO,  50 mg PO or

Alternate oral antihistamine to diphenhydramine:

- Cetirizine 10 mg  Loratadine 10 mg  Fexofenadine  60mg or  180mg

**Methylprednisolone:**  125 mg IVP  40 mg IVP or other \_\_\_\_\_ mg IVP

Famotidine:  20 mg PO,  40 mg PO,  20 mg IVP,  40 mg IVP

Pre-medicate with other: \_\_\_\_\_

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**5. DOSE:**

Rituxan® (rituximab) 1000mg IV per 500ml Sodium Chloride 0.9% to infuse per protocol OR

Other Dose: \_\_\_\_\_

**6. FREQUENCY:**

Infuse at 0 week and 2 weeks **every 4 months (16 weeks)** OR

Infuse at 0 week and 2 weeks **every 6 months (24 weeks)**

**Special orders:** \_\_\_\_\_

***If adverse drug reaction occurs, utilize the ADVERSE DRUG REACTION GUIDELINES***

**7. Physician's Signature:** \_\_\_\_\_ / \_\_\_\_\_ Date: \_\_\_\_\_

No Stamp Signatures

(Dispense as written)

(Substitution permitted)

Printed Physician's Name with Credentials: \_\_\_\_\_ NPI: \_\_\_\_\_

**8. Fax updated supporting clinical MD notes with each order renewal or change in orders**

Infusion order forms and Adverse Drug Reaction Guidelines are available at [www.palmettoinfusion.com](http://www.palmettoinfusion.com)



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## Guidelines for Prescribing Rituxan® (rituximab) (Required documentation with all initial referrals)

Patient Name: \_\_\_\_\_ Referral Date: \_\_\_\_\_

\_\_\_ Include signed and completed **Plan of Treatment**. (MD must complete sections 1-8)  
(Infusion order forms & Standard Adverse Reactions orders are available at [www.palmettoinfusion.com](http://www.palmettoinfusion.com) under Agency/MD tab)

\_\_\_ Include patient demographic information and insurance information. (Copy of insurance cards if available)

\_\_\_ **Supporting clinical MD notes to include any past tried and/or failed therapies, intolerance, outcomes or contraindications to conventional therapy. Include any lab results and/or tests to support diagnosis.**  
• Rheumatoid Arthritis (RA) Rituxan® (rituximab) in combination with methotrexate is indicated for the treatment of adult patients with moderately- to severely- active rheumatoid arthritis who have had an inadequate response to one or more TNF antagonist therapies. If patient is unable to take methotrexate, then MD must include supporting documentation as to reason/rational.

\_\_\_ If patient is switching biological therapies, then MD must specify wash-out period prior to starting Rituxan® as specified of \_\_\_\_\_ weeks. Last known biological therapy: \_\_\_\_\_ and last date received: \_\_\_\_\_. (Include copy of last Rituxan® infusion record if available and currently on therapy)

\_\_\_ Other as requested: \_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

### Pre-Screening: (Hepatitis screening results must be available prior to start of therapy and within last 12 months)

\_\_\_ **Required Hepatitis screening to include: Hepatitis B Surface Antigen (HBsAg) and Total Hepatitis B Core Antibody (anti- HBc)**  
\*Rituxan® is contraindicated in patient with active HBV. Patients who are negative for surface antigen HBsAg (-) and positive for HB core antibody HBcAB (+) or positive for surface antigen HBsAg (+), should consult liver disease experts before starting and during treatment.

\*\* Warnings/Precautions: • **Hepatitis B Virus Reactivation**- Screen all patients for HBV infection by measuring HBsAg and anti-HBc (antibodies) before initiating treatment with Rituxan®. For patients who show evidence of prior hepatitis B infection (HBsAg positive or HBsAg negative but anti-HBc positive), consult with physicians with expertise in managing hepatitis B regarding monitoring and consideration for HBV antiviral therapy before and/or during Rituxan® treatment. HBV reactivation has been reported up to 24 months following completion of Rituxan® • Glucocorticoids administered methylprednisolone IV premed or its equivalent 30 minutes prior to each infusion are recommended to reduce the incidence and severity of infusion reactions. • Subsequent courses should be administered every 24 weeks or based on clinical evaluation, but not sooner than every 16 weeks. Evaluation of immunizations should be completed prior to and live vaccines should not be given before or concurrently. **Serious Infections:** including fatal, bacterial, fungal, and new or reactivated viral infections can occur during and following the completion of Rituxan®. See full prescribing information.

Palmetto Infusion Services will complete insurance verification and submit all required clinical documentation to the patient's insurance company for eligibility. Our office will notify you if any further information is required. We will review financial responsibility with the patient and refer them to any available Co-pay assistance as required. Thank you for the referral.

Please fax all information to 1-866-872-8920 or call 1-800-809-1265 for assistance.