

**PATIENT INFORMATION**

Patient Name: \_\_\_\_\_ DOB: \_\_\_\_\_ Phone: \_\_\_\_\_ Sex: M / F Ht: \_\_\_\_\_ Wt: \_\_\_\_\_ lbs / kg  
Primary Language: \_\_\_\_\_ Allergies: \_\_\_\_\_  
Patient Preferred Location: \_\_\_\_\_

<ICD 10 CODE REQUIRED>

**DIAGNOSIS & CLINICAL INFORMATION**

**ICD 10 Code**

- M31.30 Granulomatosis w/ Polyangitis (Wegener's Granulomatosis GPA)
- M31.7 Microscopic Polyangitis
- Other: \_\_\_\_\_

**REQUIRED: Demographics & Most Recent: H&P, clinical notes, & medication list. Supporting clinical notes to include any past tried and/or failed therapies, intolerance, outcomes, or contraindications to conventional therapy.**  
**LAB RESULTS: Include Negative Hepatitis B within 3 years.**

**PRESCRIPTION**

**Pre-Medications**

Acetaminophen: 650 mg PO  
Diphenhydramine: 25 mg IVP  
Methylprednisolone: 125 mg SIVP  
Other: \_\_\_\_\_

**Ruxience (rituximab-pvvr)**

Infuse in 250-550 mL of 0.9% Sodium Chloride.

**Dose: (SELECT ONE)**

- IV: Infuse \_\_\_\_\_ mg
- IV: infuse 375 mg/m<sup>2</sup> – Required  Height: \_\_\_\_\_, Weight: \_\_\_\_\_ lbs or \_\_\_\_\_ kg

**Frequency: (SELECT ONE)**

- Once weekly for one year
- Other: \_\_\_\_\_ for one year

In the event of an adverse reaction occurring at a Medix Infusion suite, utilize the Medix Infusion adverse reaction protocol.

Post Treatment Observations: The patient is observed for 60 minutes following the first infusion.

Comments: \_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

**PRESCRIBER INFORMATION**

Prescriber Name: \_\_\_\_\_ **Signature:** \_\_\_\_\_  
Date: \_\_\_\_\_ NPI #: \_\_\_\_\_ Specialty: \_\_\_\_\_  
Supervising Physician: \_\_\_\_\_ (If Applicable)  
Address: \_\_\_\_\_ City: \_\_\_\_\_ State: \_\_\_\_\_ Zip: \_\_\_\_\_  
Contact Name: \_\_\_\_\_ Phone: \_\_\_\_\_ Fax: \_\_\_\_\_ Email: \_\_\_\_\_