

## 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 (PROVIDE COMPLETE CODE)**

M05. \_\_\_\_\_ Rheumatoid Arthritis, w/Rheumatoid Factor

M06. \_\_\_\_\_ Rheumatoid Arthritis, w/o Rheumatoid Factor

L40.5 \_\_\_\_\_ Psoriatic Arthropathy

M45 \_\_\_\_\_ Ankylosing Spondylitis

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

**REQUIRED:** Demographics & Most Recent: H&P, clinical notes, and 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 & Negative TB within 12 months. If the patient is unable to take methotrexate, then provider must include supporting documentation as to reason/rational.

## PRESCRIPTION\*

**Pre-Medications**

- Acetaminophen: 650 mg PO
- Cetirizine: 10 mg PO
- Diphenhydramine: 25 mg PO
- Diphenhydramine: 25 mg IVP
- Famotidine: 20 mg PO
- Methylprednisolone: 125 mg SIVP
- Other: \_\_\_\_\_

**Lab Orders+**

Required: Negative TB, annually

**+ Medix Infusion will draw maintenance labs unless otherwise directed by Referring Provider.**

**SIMPONI ARIA (golimumab)**

**Loading Dose**

**IV:** Infuse 2 mg/kg in 100 mL of 0.9% Sodium Chloride over at least 30 minutes via pump using 0.2-micron filter at weeks 0 and 4

**Maintenance Dose**

**IV:** Infuse 2 mg/kg in 100 mL of 0.9% Sodium Chloride over at least 30 minutes via pump using 0.2-micron filter every 8 weeks for one year

Is the patient on any other disease modifying therapy?      **Yes**      **No**

If yes, please note therapy and last dose: \_\_\_\_\_

**Post Treatment Observations:** The patient is observed for 30 minutes following the first administration.

**Adverse Events:** In the event of an adverse reaction occurring at a Medix Infusion suite, utilize the Medix Infusion adverse reactions protocol.

**Comments:**

\_\_\_\_\_  
\_\_\_\_\_

## PRESCRIBER INFORMATION

Prescriber Name: \_\_\_\_\_ Signature: \_\_\_\_\_

Date: \_\_\_\_\_ NPI #: \_\_\_\_\_ Specialty: \_\_\_\_\_

Supervising Physician: \_\_\_\_\_ (If Applicable)

Address: \_\_\_\_\_ City: \_\_\_\_\_ State: \_\_\_\_\_ Zip: \_\_\_\_\_

Contact Name: \_\_\_\_\_ Phone: \_\_\_\_\_ Fax: \_\_\_\_\_ Email: \_\_\_\_\_