



Patient Name: _____ DOB: ____/____/____ Date of Last Infusion: ____/____/____
Height _____ Weight _____

Infusion Location: (state and Site) _____

Simponi Aria® (golimumab) Infusion Orders

Diagnosis (please provide ICD-10 code in space provided):

| | |
|---------------------------------------|---|
| ____ Psoriatic Arthritis (ICD-10) | ____ Ankylosing Spondylitis (ICD-10) |
| ____ Rheumatoid Arthritis (ICD-10) | ____ Other: _____ (ICD-10) |

Nursing Orders:

- Hold infusion and notify provider for:
 - Abnormal vital signs, Fever, neurological changes, signs/symptoms of illness/active infection
 - Planned/recent surgical procedures or recent live vaccinations
- If an infusion-related reaction occurs, stop infusion and initiate Hypersensitivity Reaction Management Protocol as clinically indicated.

| | | |
|--|------------------------------|--------------------------------------|
| Lab Orders: | <input type="checkbox"/> CMP | <input type="checkbox"/> Other _____ |
| <input type="checkbox"/> CBC with diff | <input type="checkbox"/> LFT | <input type="checkbox"/> Other _____ |

Administer golimumab 2mg/kg x (current weight) _____ kg = _____ mg
in 100 mL 0.9% sodium chloride. Administer using an in-line, sterile, non-pyrogenic low-protein binding **filter** (pore size 0.22 micron or less) over a period of 30 minutes.

Frequency (chose one):

| | | |
|--|--|--|
| <input type="checkbox"/> On weeks 0, 4, then every 8 weeks | <input type="checkbox"/> Every 8 weeks | <input type="checkbox"/> Every _____ weeks |
|--|--|--|

Additional Orders:

Provider name (print) _____ Date: _____

Provider signature: _____ Time: _____

Reviewed 6/29/2022. Order valid for one year unless otherwise indicated. IV solutions/diluents may be substituted as allowed per manufacturer's

instructions as necessitated by product availability.