

## Physician's Orders GOLIMUMAB (SIMPONI ARIA), IV PIGGYBACK - ADULT, OUTPATIENT, COREWELL HEALTH INFUSION CENTER Page 1 to 2

Patient Name		
DOB		
MRN		
Physician		
CSN		

☐ Interval: <b>INDU</b>		maintenance treatment	begins on day 84)			
Anticipated Infusion Da	ite ICD 10 Code with De	scription				
Height	(cm) Weight(kg) Allergies					
Site of Service						
☐ CH Gerber	☐ CH Lemmen Holton (GR)	☐ CH Pennock	☐ CH United Memorial			
☐ CH Helen DeVos (GR)	☐ CH Ludington	☐ CH Reed City	☐ CH Zeeland			
☐ CH Blodgett (GR)						
Provider Specialty  ☐ Allergy/Immunology	□ Infactious Disease	□ OB/GYN	☐ Rheumatology			
☐ Cardiology	☐ Internal Med/Family Practice	☐ Other	☐ Surgery			
☐ Gastroenterology	☐ Nephrology	☐ Otolaryngology	• •			
☐ Genetics	□ Neurology	☐ Pulmonary	□ Wound Care			
A in-tot D	-4-	•				
Appointment Reque	SIS					
Status: Futur Infusion and	pointment Request re, Expected: S, Expires: S+365, Sched. Tolerance possible labs  nd Special Instructions	e: Schedule appointment at mo	ost 3 days before or at most 3 days after,			
<u></u>						
	TY PARAMETERS AND SPECIAL INSTI B (SIMPONI ARIA):	RUCTIONS 4				
An FDA-approved patient medication guide, which is available with the product information and as follows, should be dispensed with this medication						
Https://www.	Https://www.accessdata.fda.gov/drugsatfda_docs/label/2017/125433s019lbl.pdf					
Treatment wi infections.	Treatment with SIMPONI ARIA should not be initiated in patients with an active infection, including clinically important localized infections.					
	Tuberculosis surveillance and maintenance: Screen and treat latent infection prior to starting therapy.  Hepatitis B surveillance and maintenance: Screen prior to initiating therapy. Refer to specialist as warranted by serology.					
risk assessm active infecti splenomegal	TB skin test, hepatitis B surface antigen (HBsAg) test, liver function test (LFT), complete blood count (CBC), up-to-date vaccinations, risk assessment for cancer, and pregnancy testing. Monitor for signs of tuberculosis throughout therapy. Do not initiate therapy if active infection is present. Monitor closely for signs and symptoms of infection. Monitor for signs/symptoms of malignancy (eg, splenomegaly, hepatomegaly, abdominal pain, persistent fever, night sweats, weight loss). Identify history of latex or polysorbate 80 allergy; some dosage containers may contain these agents. Monitor LFTs, CBC at regular intervals. Assess results of laboratory tests					

+

(PDD) at regular intervals during treatment.



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Health <sup>™</sup>

### GOLIMUMAB (SIMPONI ARIA), IV PIGGYBACK - ADULT, OUTPATIENT, **COREWELL HEALTH INFUSION CENTER** Page 2 to 2

Patient Name
DOB
MRN
Physician
CSN

(CON	TINU	ED,
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Verify all INDUCTION/LOADING DOSES given prior to start of MAINTENANCE DOSES	ONC SAFETY PARAMETERS AND SPECIAL INSTRUCTIONS 6	
	Verify all INDUCTION/LOADING DOSES given prior to start of MAINTENANCE DOSES	

- ONC SAFETY PARAMETERS AND SPECIAL INSTRUCTIONS 4 HEPATITIS B VIRUS SURVEILLANCE AND MAINTENANCE RECOMMENDATIONS: Screen prior to treatment. Refer to specialist as warranted by serology.
- ONC SAFETY PARAMETERS AND SPECIAL INSTRUCTIONS 5 TUBERCULOSIS SURVEILLANCE AND MANAGEMENT RECOMMENDATIONS: Screen prior to treatment. Treat latent infection prior to starting therapy

_abs					
_				Interval	Duration
~		mplete Blood Count w/Differential May Initiate IV Catheter Patency Adult Protoco Status: Future, Expected: S, Expires:		t, Blood, Blood, Ve	enous
	H	epatic Function Panel (Liver Panel) Status: Future, Expected: S, Expires: S+365,	URGENT, Clinic Collect, Blood	l, Blood, Venous	
_	H	epatitis B Surface Antigen Level Status: Future, Expected: S, Expires: S+365,	URGENT, Clinic Collect, Blood	d, Blood, Venous	
·	И	epatitis B Core Total Antibody Level Status: Future, Expected: S, Expires: S+365,	URGENT, Clinic Collect, Blood	d, Blood, Venous	
<u>~</u> ]		rrange For Patient To Have Id Tb Skin T nerapy Or Annually	est Administered And Rea	d Or Serum Tb S	Screening Lab Prior To
	(	ONC PROVIDER REMINDER 28 Arrange for patient to have intradermal TB ski annually.	n test (tuberculin PPD) screeni	Once ng performed and r	1 treatment read prior to initiating therapy and
	-	FB Screen (Quantiferon Gold) Status: Future, Expected: S, Expires: S+365,	URGENT, Clinic Collect, Blood	Once d, Blood, Venous	1 treatment
Additio	nal	Lab Orders			
			Interval		Duration
		Labs:	☐ Every ☐ Once	days	□ Until dat □ 1 year □ # of





# GOLIMUMAB (SIMPONI ARIA), IV PIGGYBACK - ADULT, OUTPATIENT, COREWELL HEALTH INFUSION CENTER (CONTINUED) Page 3 to 2

**Patient Name** DOB MRN Physician

CSN

**Nursing Orders** 

Interval

Duration

Until discontinued

Hypersensitivity Reaction Adult Oncology Protocol

Routine, Until discontinued Starting when released for 24 hours HYPERSENSITIVITY REACTIONS:

Discontinue the medication infusion immediately.

Activate emergency response for severe or rapidly progressing symptoms. Where available consider calling RAP and have crash cart available. Call 911 or code team (if applicable) as needed for an absence of pulse and respirations. Refer to site specific emergency response policy.

Stay with patient until symptoms have resolved.

Initiate/Continue Oxygen to maintain SpO2 greater than 90% and discontinue Oxygen Therapy to maintain SpO2 above 90%

For severe or rapidly progressing hypersensitivity reaction symptoms, monitor vital signs and pulse oximeter readings every 2 to 5 minutes until the patient is stable and symptoms resolve.

Document medication infusing and approximate dose received at time of reaction in the patient medical record. Document allergy to medication attributed with causing reaction in patient medical record. Complete Adverse Drug Reaction form per Pharmacy Clinical Policy.

May Initiate IV Catheter Patency Adult Protocol.

### **Treatment Parameters**

ONC MONITORING AND HOLD PARAMETERS 3

May proceed with treatment if hepatitis B core antibody and surface antigen labs have been resulted prior to the first dose, and the results are negative.

ONC MONITORING AND HOLD PARAMETERS 4

May proceed with treatment if tuberculosis screening test with either TB Screen blood test (QuantiFERON® Gold Plus) or TB skin test have been resulted prior to first dose and the results are negative.

### Medications

golimumab (SIMPONI ARIA) 2 mg/kg in sodium chloride 0.9 % 100 mL IVPB

2 mg/kg, Intravenous, Administer over 30 Minutes, Once, Starting S, For 1 Dose

Infuse over 30 minutes. Do not infuse in the same line with other medications.

Telephone order/Verbal order documented and read-back completed. Practitioner's initials \_

NOTE: Unless Order is written DAW (dispense as written), medication may be supplied which is a generic equivalent by nonproprietary name.

TRANSCRIBED:		VALIDATED:		ORDERED:			
TIME	DATE	TIME	DATE	TIME	DATE	Pager#	
			R.N.		Physic	ian	Physician
	Sign		Sign		P	rint	Sign

EPIC VERSION DATE: 09/12/20

