

## Physician's Orders TOCILIZUMAB -PEDIATRIC, OUTPATIENT, INFUSION CENTER

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Patient Name
DOB
MRN
Physician
FIN

Defaults for orders not oth Interval: Every 7 days Interval: Every 14 days Interval: Every 21 days	· S		
Duration:  Until date:  1 year  multing freatment			
Anticipated Infusion Date_	ICD 10 Code with Des	cription	
Height(cm	) Weight(kg) Allergies_		
<b>Provider Specialty</b>			
□ Allergy/Immunology	☐ Infectious Disease	☐ OB/GYN	□ Rheumatology
☐ Cardiology	☐ Internal Med/Family Practice	☐ Other	☐ Surgery
☐ Gastroenterology	☐ Nephrology	□ Otolaryngology	☐ Urology
☐ Genetics	☐ Neurology	☐ Pulmonary	☐ Wound Care
Site of Service		□ SH Pennock	CI I United Mensonial
☐ SH Gerber	<ul><li>☐ SH Lemmen Holton (GR)</li><li>☐ SH Ludington</li></ul>		<ul><li>☐ SH United Memorial</li><li>☐ SH Zeeland</li></ul>
☐ SH Helen DeVos (GR)	☐ SH Ludington	☐ SH Reed City	□ S⊓ Zeelaliu
Provider Reminder  ONC PROVIDER I  Premedication is not r reactions requiring tre	ted: S, Expires: S+365, Sched. Tolerance: Sche	red in patients who were premec	dicated. In clinical studies,
Select Tocilizumab M	ledication Instructions Based On R	oute	
ONC NURSING COM	MMUNICATION 1		
Tocilizumab (INTRAVENOU	S)		
- Do not administer if the	solution is discolored or if foreign particulate ma	atter is present.	
- Place intermittent infusi	•		
anaphylaxis every 15 mir - Notify physician, NP or bronchospasm.	obtain heart rate, respiratory rate, blood pressur- nutes during infusion through 30 minutes after in PA-C and stop infusion immediately if patient ha PA-C if greater than 20% decrease in systolic o	fusion. as itching, hives, swelling, fever,	•
hypersensitivity or anaph	liphenhydramine/Epi-pen available (as appropria ylactic reactions may occur during and after infu tions, and importance of seeking medical care.		

- At the end of IV infusion, flush secondary line with 0.9% Sodium Chloride.



## TOCILIZUMAB PEDIATRIC, OUTPATIENT, INFUSION CENTER (CONTINUED)

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## ☐ ONC NURSING COMMUNICATION 2

Tocilizumab (SUBCUTANEOUS)

- Do not administer if the solution is discolored or if foreign particulate matter is present.
- For subcutaneous administration, allow medication to reach room temperature prior to use. Rotate injection sites; avoid injecting into moles, scars, or tender, bruised, red, or hard skin.
- For subcutaneous route, obtain heart rate, respiratory rate, blood pressure, and pulse oximetry and assess for symptoms of anaphylaxis every 15 minutes through 30 minutes after injection.
- Notify physician, NP or PA-C if patient has itching, hives, swelling, fever, rigors, dyspnea, cough, or bronchospasm.
- Notify physician, NP or PA-C if greater than 20% decrease in systolic or diastolic blood pressure.
- Verify that patient has diphenhydramine/Epi-pen available (as appropriate) for immediate home use. Advise patient that severe hypersensitivity or anaphylactic reactions may occur during and after injection. Inform patients of signs and symptoms of anaphylaxis and hypersensitivity reactions, and importance of seeking medical care.

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			Interval	Duration
		Complete Blood Count w/Differential	□ Every days	□ Until date:
		STAT, Starting S, For 1 Occurrences, Blood, Venous	□ Once	□ 1 year
				□# of Treatments
-		Comprehensive Metabolic Panel (CMP)	- Firemi devis	
		STAT, Starting S, For 1 Occurrences, Blood, Venous	□ Every days □ Once	□ Until date: □ 1 year
		, , , ,	□ Office	□ # of Treatments
		Sedimentation rate	□ Every days	□ Until date:
		STAT, Starting S, For 1 Occurrences, Blood, Venous	□ Once	□ 1 year
				□# of Treatments
		Lipid Panel	□ Every days	□ Until date:
		STAT, Starting S, For 1 Occurrences, Blood, Venous	□ Once	□ 1 year
				□# of Treatments
٨٨	ditiona	l Lab Orders		
Au	uitiona	i Lab Orders	Interval	Duration
			_ F	
			□ Everydays □ Once	□ Until date: □ 1 year
			□ Office	□ # of Treatments
			☐ Everydays	☐ Until date:
ш			Once	□ 1 year
			_ C.1.65	□ # of Treatments
Pre	a-Medic	cations		
Pre	e-Medic	cations		
		cations edication with dose:		

☐ Pre-medication with dose:

## TOCILIZUMAB PEDIATRIC, OUTPATIENT, INFUSION CENTER (CONTINUED) Page 3 of 3

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Medicatio	ons						
Select Tocilizumab For Intravenous Or Subcutaneous Administration							
	tocilizumab (ACTEMRA) in sodium chloride 0.9 % IVPB						
	Dose:						
	□ 4 mg/kg						
	□ 8 mg/kg						
	□ 10 mg/kg						
	□ 12 mg/kg						
	Intravenous, Administer over 60 Minutes, Once, Starting S, For 1 Doses						
	Maximum dose is 800 mg.						
	Protect from light. Do NOT shake.						
	Use a dedicated IV line. Do not administer IV push or IV bolus.						
	Do not use if opaque particles or discoloration is visible.						
	tocilizumab (ACTEMRA) subcutaneous prefilled syringe 162 mg						
	162 mg, Subcutaneous, Once, Starting S, For 1 Doses						
	Allow to reach room temperature prior to use. Rotate injection sites.						

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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 #
	Sign		R.N. Sign		Physician Print	Physician

**EPIC VERSION DATE:** 03/27/20