

## **Edward-Elmhurst Cancer Centers**

120 Spalding Drive; Suite 111; Naperville, IL 60540 Phone: 630/646-2273 Fax: 630/548-6617

24600 W. 127<sup>th</sup> Street; Plainfield, IL 60585 Phone: 630/646-2273 Fax: 630/548-6617 177 E. Brush Hill Road; Elmhurst, IL 60126 Phone: 630/646-2273 Fax: 331/221-3887

## Actemra (TOCILIZUMAB) Standing Order

Patient Name: DOB:					
***Please include current history and physical and any recent labs/tests, if applicable.***					
*PLEASE ATTACH COPY OF INSURANCE CARD WITH THIS ORDER*					
Pre-Authorization # or Call Reference #:  (Ordering Physician Office is Responsible to Obtain Authorization/Referral)					
Contact Name and Phone Number of Insurance Company:					
If you have any questions regarding pre-authorizations, please contact (630) 527-3788 and ask for the billing department.					
Allergies:					
Diagnosis (ICD 10 Required):					
Weight: Lbs: Kg:					
Actemra is indicated for treatment of adult patients with moderately to severely active RA who had an inadequate response to 1 or more TNF antagonist therapies. Recommended starting dose is 4mg/kg every 4 weeks, increasing to 8mg/kg every 4 weeks based on clinical response.					
Pre-administration:  Current TB skin test or chest x-ray ANC, Platelet count, Liver enzymes (ALT, AST) (Not recommended to start Actemra if ANC < 2000/mm3, platelets < 100,000/mm3 and liver enzymes > 1.5 times upper limit of normal) Pre-Infusion assessment to include BP, temp, pulse per unit protocol Post-infusion assessment to include BP, temp, pulse per unit protocol					
Labs:  1. Please draw the following labs (please check):  CBC w/platelets and Auto Differential Liver Enzymes Other:  Desired Frequency (please check):  Every 4 weeks done One week prior to infusion Every 8 weeks done One week prior to infusion Other: Other:					
2. Please draw the following labs (please check):  Lipid Panel					

	-						
	Desired Frequency (please check):						
	One week prior to fir	st infusion,	usion, then 4 weeks after treatment initiation, then every				
	6 months						
		st infusion,	then 8 weeks after treatr	ment initiation, then every			
	6 months						
		ent initiation	n, then every 6 months				
	U Other:						
۸ ۵۰	ministration:						
1.		ka for	mg of Actemra eve	ony 1 wooks			
١.	Maximum dose is 80		IIIg of Acternia eve	ary 4 weeks.			
	Maximum dose is ou	onig					
2.	Infuse 100ml Actemra solution	over 60 mi	nutes. Do not administe	r as bolus or push.			
3.	Do not administer Actemra dur	ing an activ	e infection, including loc	alized infections.			
4.	Interrupt Actemra:						
⊣.	<u>-</u>	develons i	ncluding localized infecti	ons			
	ANC 500-1000	dovolopo, ii	noidaing localized infoot	0110			
	Platelet count 50,000	0-100.000					
			ter modification of DMA	RD dose (4mg/kg dose)			
	Liver Enzymes > 3 to	5X ULN		, ,			
5.	Discontinue Actemra:						
	ANC < 500						
	Platelets < 50,000						
	Liver enzymes > 5X	ULN					
6	Reduce Actemra dose from 8mg/kg to	o 1 m a/l.a.					
0.			ter modification of DMA	RD dose			
0.			ter modification of DMA	RD dose			
		3X ŪLN af					
7.	Liver enzymes > 1 to	o 3X ULN at					
7.	Liver enzymes > 1 to  Do not administer Actemra concomita	o 3X ULN at					
7.	Liver enzymes > 1 to  Do not administer Actemra concomita in combination with other biologic DM	o 3X ULN at					
7.	Liver enzymes > 1 to  Do not administer Actemra concomita in combination with other biologic DM  Infusion reactions may include:	o 3X ŪLÑ af antly in the s IARDS.	same line with other drug	gs. Avoid use of Actemra			
7.	Liver enzymes > 1 to  Do not administer Actemra concomitation combination with other biologic DM  Infusion reactions may include:  *Headache	o 3X ŪLÑ af antly in the s IARDS. *Hives/į	same line with other drug	gs. Avoid use of Actemra  *Hypertension			
7.	Liver enzymes > 1 to  Do not administer Actemra concomitation combination with other biologic DM  Infusion reactions may include:  *Headache *Shortness of breath	o 3X ŪLÑ af antly in the s IARDS. *Hives/j *Backad	same line with other drug oruritis che	gs. Avoid use of Actemra  *Hypertension *Dizziness			
7. 8.	Liver enzymes > 1 to  Do not administer Actemra concomitation combination with other biologic DM  Infusion reactions may include:     *Headache     *Shortness of breath     *Nausea/vomiting	*Hives/p	same line with other drug oruritis che tightness	*Hypertension *Dizziness *Anaphylaxis			
7. 8. 9.	Liver enzymes > 1 to  Do not administer Actemra concomitation combination with other biologic DM  Infusion reactions may include:     *Headache     *Shortness of breath     *Nausea/vomiting  In the event of a hypersensitivity re	*Hives/p *Backad *Chest	same line with other drug oruritis che tightness ring the infusion of this	*Hypertension *Dizziness *Anaphylaxis *medication, we will			
<ul><li>7.</li><li>8.</li><li>9.</li></ul>	Liver enzymes > 1 to  Do not administer Actemra concomits in combination with other biologic DM  Infusion reactions may include:	*Hives/p *Backac *Chest	same line with other drug oruritis che tightness ring the infusion of this ed nurse practitioner w	*Hypertension *Dizziness *Anaphylaxis *medication, we will			
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<ul> <li>Methylprednisolone 100mg IV Push</li> </ul>	IV Push	Yes 🗌 No 🗌		
12. Patient may be discharged post-infusion if stable. No monitoring time required.				
Physician Signature:	Date:			
Ordering Physician NPI:	Edward Hospital NP	PI: 1427069632		
	Elmhurst Hospital N	NPI: 1548306343		
Physician Name (Please Print)	Office Phone	Fax Number		

Revision/Review Date: 07/01/2021