



TOCILIZUMAB

(Actemra)

Outpatient Infusion

Order Form

Patient Name	
DOB	
Address	
Phone	

Order Status New Order Renewal Dose or Frequency Change

Allergies:	Weight:	Height:
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Diagnosis	<input type="checkbox"/> Rheumatoid arthritis	<input type="checkbox"/> Cytokine release syndrome	Diagnosis Code:
	<input type="checkbox"/> Systemic sclerosis associated interstitial lung disease	<input type="checkbox"/> Giant cell arteritis <input type="checkbox"/> Other:	

Required Information	Negative Quantiferon TB, T-spot or chest x-ray (no active disease)	Date:
	Diagnostic Hepatitis B panel	Date:
	Coccidioides Screen/Panel	Date:
	CBC and CMP	Date:

Required Documentation

- H&P or progress note supporting diagnosis and medication history
- RA: Positive RF or anti-CCP or RF, Anti-CCP and CRP and/or ESR and inadequate response to methotrexate or another convention synthetic drug or inadequate response to NSAIDs and/or intra-articular glucocorticoids
- Giant cell arteritis: Temporal artery biopsy or cross-sectional imaging or acute-phase reactant elevation and/or high CRP
- SSc-ILD: HRCT study of chest
- Cytokine release syndrome: CAR T cell-induced CRS or refractory CRS related to blinatumomab therapy
- Recent labs (as above) and/or diagnostic test results

Labs	<input type="checkbox"/> Hepatic Function panel every 3 months	<input type="checkbox"/> Other:
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Pre-Medications

- | | | | | |
|--|--------------------------|-----------------------------|--|------------------------------|
| <input type="checkbox"/> Diphenhydramine | <input type="radio"/> PO | <input type="radio"/> 25 mg | <input type="checkbox"/> Acetaminophen | <input type="radio"/> 325 mg |
| | <input type="radio"/> IV | <input type="radio"/> 50 mg | | <input type="radio"/> 650 mg |
| <input type="checkbox"/> Other: | | | | |

Actemra (tocilizumab) IV Medication Order

- Actemra (tocilizumab) IV 4 mg/kg every ____ weeks
- Actemra (tocilizumab) IV 6 mg/kg every ____ weeks
- Actemra (tocilizumab) IV 8 mg/kg every ____ weeks
- Other:

Infusion Reaction Medications

Hypersensitivity Reaction Protocol will be utilized unless otherwise specified

Provider (print name):	Date:
Provider Signature:	NPI:
Office Phone:	Office Fax: