

Referral Status:	MRN:
<input type="checkbox"/> New referral	<input type="checkbox"/> Order change
<input type="checkbox"/> Order Renewal	
Patient preferred clinic:	

# Soliris® (eculizumab) Standard Plan of Treatment for aHUS

Rev 4.29.25

## PATIENT DEMOGRAPHICS:

Patient Name:	
Patient's Phone:	Address:
Date of Birth:	City, State, Zip:
Height in inches:	Weight: LB or KG
Gender:	Allergies:
	<input type="checkbox"/> See list <input type="checkbox"/> NKDA

## DIAGNOSIS: (PLEASE COMPLETE 2<sup>ND</sup> AND 3<sup>RD</sup> DIGITS TO COMPLETE ICD 10 FOR BILLING)

D58.8 - Other specified hereditary hemolytic anemias	D59.30 - Atypical Hemolytic Uremic Syndrome (aHUS)
D59.8 - Other acquired hemolytic anemias	D59.32 - Hereditary hemolytic - uremic syndrome
D59.39 - Other hemolytic- uremic syndrome	- Other:
D59.4 - Other non autoimmune hemolytic anemias (including microangiopathic hemolytic anemia)	

## REQUESTED DOCUMENTATION:

## PREVIOUS ADMINISTRATION: HAS THIS PATIENT TAKEN THIS MEDICATION BEFORE?

1	Insurance information	IF NO:	IF YES:
2	History & Physical/Tried and failed therapies	PLEASE STATE	LAST INFUSION DATE:
3	Full medication list	REQUIRED WASHOUT	NEXT INFUSION DATE:
4	<b>REQUIRED:</b> Documentation of meningococcal vaccine (MenACWY AND MenB) at least 2 weeks prior to start of therapy	FROM PREVIOUS THERAPY:	<b>IF ORDER CHANGE:</b>
			<b>Continue current order until insurance approved</b>

## MEDICATION ORDERS:

NOTE: Patient may be ineligible to receive eculizumab if receiving antibiotics for active infectious process, antifungal therapy, active fever and/or suspected infection, presents with any symptoms of meningococcal infections, and/or surgery.

## PREMEDICATION TO BE ADMINISTERED 30 MINUTES PRIOR TO ADMINISTRATION AS SELECTED

\*FDA labeling does not suggest any premedication prior to infusion

IV	Diphenhydramine	25mg	50mg		PO	Acetaminophen	325mg	500mg	650mg	1000mg
	Methylprednisolone	40mg	125mg	Other:		Famotidine	20mg	40mg		
	Famotidine	20mg	40 mg			Diphenhydramine	25mg	50mg		
	Other:					Fexofenadine	60mg	180mg		
						Cetirizine	10mg			
					Loratadine	10mg				
					Other:					

## MEDICATION:

Soliris® (eculizumab) IV given over 35 minutes diluted in NS according to FDA labeling instructions  
**If the infusion is slowed, the total infusion time should not exceed 2 hours.**

\*Follow each infusion with a 1 hour post infusion monitoring\*

## SPECIAL/OTHER LAB ORDERS:

<input checked="" type="checkbox"/>	Refills x 12 months unless noted otherwise here:
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## FREQUENCY/DOSE:

<input type="checkbox"/>	Induction: 900mg/180ml NS IV weekly for 4 weeks
<input type="checkbox"/>	Maintenance (to begin at week 5 if receiving induction): 1200mg/240ml NS IV every 2 weeks
<input type="checkbox"/>	Other:

**Prescriber must be enrolled in the Soliris (REMS) program, at 1 888 765 4747 or at [www.solirisrems.com](http://www.solirisrems.com).**

## LINE USE/CARE ORDERS:

- Start PIV/Access CVC
- Flush device per facility standard flushing procedure
- Provide nursing care per Palmetto Infusion Nursing Procedures and post procedure observation if indicated

## ADVERSE REACTION & ANAPHYLAXIS ORDERS:

Administer acute infusion and anaphylaxis medications per Palmetto Infusion standing adverse reaction orders, which can be found at our website.

## PRESCRIBER INFORMATION:

PROVIDER NAME:	PHONE:
ADDRESS:	FAX:
CITY, STATE, ZIP:	NPI:

## PRESCRIBER SIGNATURE: (No stamp signatures)

## DATE:

Dispense as written/Brand medically necessary	Substitution permitted