

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 Neuromyelitis Optica Spectrum Disorder

**PATIENT DEMOGRAPHICS:**

Date of Referral:	Patient's Phone:
Patient Name:	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)**

G36.0 - Neuromyelitis Optica Spectrum Disorder	- Other:
--	----------

**REQUESTED DOCUMENTATION:**

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

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

**MEDICATION:**

Soliris® (eculizumab) IV given over 35 minutes in diluted in NS per FDA labeling suggestions

**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:**

\_\_\_\_\_

\_\_\_\_\_

**FREQUENCY/DOSE:**

Induction: 900mg in 180ml NS IV given weekly for 4 weeks

Maintenance (to begin at week 5 if receiving induction): 1200mg in 240ml NS IV given once every 2 weeks

Other: \_\_\_\_\_

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

Refills x 12 months unless noted otherwise here:

**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 or scan here.


**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
---	------------------------