

# ECULIZUMAB (Soliris) Infusion Order Form

Original ordering provider must be enrolled in REMS program <https://www.ultsolrems.com/>

Weight: \_\_\_\_\_ kg Height: \_\_\_\_\_ inches [ ] New treatment  
[ ] Transfer - Start on week # \_\_\_\_\_

## **PRIOR TO FIRST DOSE:**

Verify meningococcal vaccine completed at least 14 days prior to start date.#

*Note: Do not use combined ABCWY vaccine as full immune effect delayed for 6 months.*

- Meningococcal conjugate (MenACWY) vaccine  
0.5 ml IM x 2 doses at least 8 weeks apart **Date complete** \_\_\_\_\_  
Booster 0.5 ml IM x1 every 5 years.\*
  - Meningococcal group B (MenB) vaccine  
0.5 ml IM x 2 doses at least 4 weeks apart **Date complete** \_\_\_\_\_  
Booster 0.5 ml IM once year after series complete, then every 2 to 3 years.\*
- \* Note dates booster dose due in clinic record.

# **If less than 2 weeks since full meningococcal vaccination**, verify patient is on prophylactic anti-infective for the first 2 weeks of therapy. Consider antimicrobial prophylaxis with oral antibiotics for the duration of therapy.

Antimicrobial therapy ordered if any: \_\_\_\_\_

## **PRIOR TO EACH INFUSION**

See page 2 for labs, nursing assessment, vital signs, administration and observation times

**PRE-MEDICATE** (30 minutes before infusion): [ ] loratadine 10 mg PO  
[ ] acetaminophen \_\_\_\_\_ mg PO [ ] diphenhydramine \_\_\_\_\_ mg PO  
[ ] methylprednisolone \_\_\_\_\_ mg IV [ ] diphenhydramine \_\_\_\_\_ mg IV

**DOSE** (select diagnosis): Administer within 2 days of the recommended interval.

[ ] **Myasthenia Gravis-*AChR positive*** [ ] **Neuromyelitis optica- *aquaporin-4-antibody positive***  
[ ] **Atypical hemolytic uremic syndrome**  
Eculizumab 900 mg IV weekly x 4 weeks, 1200 mg IV week 5, then continue 1200 mg IV every 2 weeks

[ ] **Paroxysmal Nocturnal Hemoglobinuria (PNH)**  
Eculizumab 600 mg IV weekly x 4 weeks, 900 mg IV week 5 then continue 900 mg IV every 2 weeks

[ ] Other: \_\_\_\_\_

**REQUIRED Prior Authorization Number:** \_\_\_\_\_ [ ] pending [ ] complete [ ] not needed\*

\*If not needed is chosen, date, time and name of person at health insurer who authorized.

Date: \_\_\_\_\_ Time: \_\_\_\_\_ Duration of authorization: \_\_\_\_\_

## **Checklist for non-RFGH credentialed providers:**

- [ ] Problem list, current medication and allergies attached
- [ ] Provider to provider communication is required. If the patient has a Primary Care Provider at RFGH, please contact that PCP. Otherwise, call (207) 474-5121 and ask to speak to hospitalist .  
Contacted provider: \_\_\_\_\_

**FAX RFGH Infusion clinic at 207-858-2404**

**Contact Infusion Clinic at 207-858-8722**

Revised 7/8/25

Provider signature \_\_\_\_\_ Date \_\_\_\_\_ time \_\_\_\_\_

*If not RFGH credentialed:*

Printed name \_\_\_\_\_ NPI # \_\_\_\_\_ (required)

**RFGH Co-signature** \_\_\_\_\_

Date \_\_\_\_\_ time: \_\_\_\_\_

Patient name \_\_\_\_\_

Date of birth \_\_\_\_\_

Patient phone number \_\_\_\_\_

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## **PRIOR TO EACH INFUSION:**

- CBC with differential, lactic dehydrogenase (LDH), serum creatinine, AST, Urinalysis – culture if indicated before each treatment
- Supply patient with the manufacturer/FDA Medication Guide and medication safety card.
- Instruct patient of signs of infusion type reaction and to immediately report chest pain, trouble breathing, or swelling of face, tongue, or throat.
- Assess for infection; delay administration for active infection.

## **INFUSION**

- Vital signs prior to infusion and every 15 minutes.
- Infuse over 35 minutes. Decrease infusion rate or discontinue for infusion reactions;
- Do not exceed a maximum 2-hour duration of infusion in adults.
- Observe patient for 1 hour after infusion.

Revised: 4/8/25

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