

**SOLIRIS / ULTOMIRIS  
REFERRAL FORM**

**Y Medical Associates**  
**Fax: 855-838-0623**  
**Phone: 800-447-7558**



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

**Patient Information**

Patient Name: \_\_\_\_\_  
 Address: \_\_\_\_\_  
 City, State, Zip: \_\_\_\_\_  
 Home Phone: \_\_\_\_\_  
 Cell Phone: \_\_\_\_\_  
 DOB: \_\_\_\_\_ Gender:  M  F

**Prescriber Information**

Prescriber Name: \_\_\_\_\_  
 Address: \_\_\_\_\_  
 City, State, Zip: \_\_\_\_\_  
 Phone: \_\_\_\_\_ Fax: \_\_\_\_\_  
 DEA: \_\_\_\_\_ NPI #: \_\_\_\_\_  
 Contact Person: \_\_\_\_\_

**INSURANCE INFORMATION (Please attach the front and back of insurance and prescription drug cards)**

Primary Insurance: \_\_\_\_\_ ID#: \_\_\_\_\_ Group: \_\_\_\_\_  
 Secondary Insurance: \_\_\_\_\_ ID#: \_\_\_\_\_ Group: \_\_\_\_\_  
 Prescription Card: \_\_\_\_\_ ID#: \_\_\_\_\_ BIN#: \_\_\_\_\_ PCN#: \_\_\_\_\_ Group: \_\_\_\_\_

**DIAGNOSIS & CLINICAL ASSESSMENT (Fill in below or attach lab work)**

New to Therapy  Currently on Therapy  Date of Last IVIG Infusion: \_\_\_\_\_  IVIG Dosing Regimen: \_\_\_\_\_

**Diagnosis:**  G70.00 Myasthenia Gravis without (acute) exacerbation  G70.01 Myasthenia Gravis with (acute) exacerbation  in crisis

D59.3 atypical Hemolytic Uremic Syndrome (aHUS)  D59.5 PNH  G36.0 Neuromyelitis Optica **Date of Diagnosis:** \_\_\_\_\_

**Weight:** \_\_\_\_\_ **Height:** \_\_\_\_\_ **Date:** \_\_\_\_\_ **Allergies:** \_\_\_\_\_ **Date of MenACWY:** \_\_\_\_\_ **Date of MenB:** \_\_\_\_\_

**Previously on PLEX treatment**  Yes  No **Date of last treatment:** \_\_\_\_\_ **Is patient AchR antibody positive?**  Yes  No

**Is the Patient Anti-Aquaporin-4 (AQP4) antibody positive?**  Yes  No **Notes/Comments:** \_\_\_\_\_

**Soliris (eculizumab)**

**Ultomiris (ravulizumab)**

**Strength**

**Directions**

Injection:  
 300mg / 30mL  
 (10mg/mL) in a  
 single-dose vial

- For treatment of Myasthenia Gravis:**
  - 900mg weekly for the first 4 weeks, followed by  1200mg for the fifth dose 1 week later, then  1200mg every 2 weeks thereafter.
- For treatment of aHUS – 18 years or older:**
  - 900mg weekly for the first 4 weeks, followed by  1200mg for the fifth dose 1 week later, then  1200mg every 2 weeks thereafter.
- For treatment of NMOSD:**
  - 900mg weekly for the first 4 weeks, followed by  1200mg for the fifth dose 1 week later, then  1200mg every 2 weeks thereafter.
- For treatment of PNH – 18 years or older:**
  - 600mg weekly for the first 4 weeks, followed by  900mg for the fifth dose 1 week later, then  900mg every 2 weeks thereafter.

Orders and Medications

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

**Strength**

**Directions**

Injection:  
 300mg/30mL  
 (10mg/mL) in a  
 single dose vial  
 300mg/3 mL  
 (100mg/mL) in a  
 single dose vial  
 1,100mg/11mL  
 (100mg/mL) in a  
 single dose vial

**\* Do not mix ULTOMIRIS 100 mg/mL (3mL and 11mL vials) and 10mg/mL (30mL vial) concentrations together.**

**\*\*When switching therapy:**  
 Ultomiris loading dose should be given at the time of next scheduled Soliris dose.

- For treatment of Myasthenia Gravis - weight based at time of treatment (patient must be at least 40kg):**
  - \_\_\_\_\_ mg as a single dose, followed by  \_\_\_\_\_ mg once every 8 weeks later starting 2 weeks after the loading dose.
- For treatment of aHUS – weight based at time of treatment:**
  - \_\_\_\_\_ mg as a single dose, followed by  \_\_\_\_\_ mg once every \_\_\_\_\_ (4 or 8) weeks later starting 2 weeks after the loading dose.
- For treatment of PNH – weight based at time of treatment:**
  - \_\_\_\_\_ mg as a single dose, followed by  \_\_\_\_\_ mg once every \_\_\_\_\_ (4 or 8) weeks later starting 2 weeks after the loading dose.

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

**Anaphylaxis Orders**

Diphenhydramine Administer 25 mg slow IV/IM may repeat x1 **Dispense:** 1 x 50 mg vial  
 Epinephrine Autoinjector  Administer 0.15mg (1:2000) IM (< 30 Kg) **Dispense:** 1 package (2 pens)  
 Administer 0.3mg (1:1000) IM (≥ 30 Kg)  
 Sodium Chloride 0.9% Use to maintain IV line, prevent or treat hypotension in case of anaphylaxis **Dispense:** QS

**IV Access Flush Order:**

NaCl 0.9% 5-10ml IV before and after infusion  
 Heparin 10 Units/ml 5ml after infusion for PICC/Midline Heparin 10 Units/ml 3ml after infusion for PIV  
 Heparin 100 Units/ml 5ml IV after infusion for PORT  
 All infusion supplies necessary to administer the medication

**Quantity**

\_\_\_\_\_

**Refills**

\_\_\_\_\_

**Other/Notes:**

\_\_\_\_\_  
 \_\_\_\_\_  
 \_\_\_\_\_

**Prescriber Signature:** \_\_\_\_\_ **DAW (Dispense as Written)**  Y  N **Date:** \_\_\_\_\_