

Ultomiris™ (ravulizumab) Standard Plan of Treatment

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 | NKDA |

DIAGNOSIS: (PLEASE COMPLETE 2ND AND 3RD DIGITS TO COMPLETE ICD 10 FOR BILLING)

| | |
|---------------------------------------------|-------------------------------------------------------|
| D59.5 - Paroxysmal nocturnal hemoglobinuria | G70.01 - Myasthenia Gravis without acute exacerbation |
| D59.3 - Hemolytic Uremic Syndrome | G70.00 - Myasthenia Gravis without acute exacerbation |
| - Other: | |

REQUESTED DOCUMENTATION:

| | | | |
|---|-----------------------------------------------------------------------------------------------------------------------|------------------------|--------------------------------------------------------|
| 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 | REQUIRED: Documentation of meningococcal vaccine (MenACWY AND MenB) at least 2 weeks prior to start of therapy | FROM PREVIOUS THERAPY: | IF ORDER CHANGE: |
| | | | Continue current order until insurance approved |

MEDICATION ORDERS:

NOTE: Patient may be ineligible to receive ravulizumab 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

| | | | | | | | | | | |
|-----------|--------------------|------|-------|--------|-----------|-----------------|-------|-------|-------|--------|
| 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:

☒ Ultomiris™ (ravulizumab) to be diluted in NS. Infused via IV per protocol.

DOSE:

☐ Dosage based on the following guidelines from the FDA package labeling.

| Patient Body Weight | Initial Dose | Maintenance Dose/Interval |
|-------------------------|--------------|---------------------------|
| 40kg to less than 60kg | 2400mg | 3000mg |
| 60kg to less than 100kg | 2700mg | 3300mg |
| 100kg or greater | 3000mg | 3600mg |

every 8 weeks

FREQUENCY:

☐ Loading dose at week 0 followed by maintenance dose at week 2 and every 8 weeks thereafter

☐ Maintenance dosing every 8 weeks

☐ Other: _____

SPECIAL/LAB ORDERS

☐ Other: _____

Follow each infusion with a 1-hour post observation.

Ultomiris™ (ravulizumab) is restricted to credentialed prescribers enrolled in the Ultomiris (REMS) program.



Refills x 12 months unless noted otherwise here:

LINE USE/CARE ORDERS:

- ☒ Start PIV/Access CVC
- ☒ Flush device per facility standard flushing procedure

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 |
|-----------------------------------------------|------------------------|