

# Ravulizumab-cwvz (Ultomiris)

**REFERRAL STATUS:**  New Referral  Dose or Frequency Change  Order Renewal

Infusion Office Preference: \_\_\_\_\_

**PATIENT INFORMATION**

Date:	Patient Name:	DOB:
<input type="checkbox"/> NKDA Allergies:	Weight (lbs / kg):	Height:
Patient Status: <input type="checkbox"/> New to Therapy <input type="checkbox"/> Continuing Therapy - Last Treatment Date:	Next Due Date:	

**PROVIDER INFORMATION**

Office Contact Name:	Office Email:
Prescribing Providers Name:	Provider NPI:
Office Address:	City: State: Zip:
Office Phone Number:	Office Fax Number:

**DIAGNOSIS AND ICD 10 CODE**

<input type="checkbox"/> Myasthenia gravis without (acute) exacerbation	ICD-10 Code: G70.00
<input type="checkbox"/> Myasthenia gravis with (acute) exacerbation	ICD-10 Code: G70.01
<input type="checkbox"/> Paroxysmal Nocturnal Hemoglobinuria (PNH)	ICD 10 Code: D59.5
<input type="checkbox"/> Neuromyelitis Optica (NMO), Aquaporin 4 Antibody Positive	ICD 10 Code: G36.0
<input type="checkbox"/> Hemolytic-uremic syndrome (aHUS)	ICD 10 Code: D59.3

**REQUIRED DOCUMENTATION/Testing**

<input type="checkbox"/> This signed order form by the provider	<input type="checkbox"/> Acetylcholine Receptor Antibody Test Results (if Myasthenia Gravis)
<input type="checkbox"/> Patient demographics AND insurance info	<input type="checkbox"/> Documentation of meningococcal vaccines
<input type="checkbox"/> Clinical/Progress notes supporting primary dx	

Is your patient enrolled in the Ultomiris-REMS program?  YES  N  
 Is the ordering PROVIDER enrolled in the Ultomiris-REMS program?  YES  N (if no, must be enrolled to start therapy)

List Tried & Failed Therapies (if Myasthenia Gravis)  
 1) \_\_\_\_\_ 2) \_\_\_\_\_

**MEDICATION ORDER**

Initial Dosing	<input type="checkbox"/> 2,400 mg IV (40k to less than 60kg) <input type="checkbox"/> 2,700 mg IV(60k to less than 100 kg) <input type="checkbox"/> 3,000 mg IV (100k or greater kg)
Maintenance Dosing	<input type="checkbox"/> 3,000 mg (40k to less than 60kg) IV every 8 weeks starting 2 weeks after initial load <input type="checkbox"/> 3,300 mg (60k to less than 100 kg) IV every 8 weeks starting 2 weeks after initial load <input type="checkbox"/> 3,600 mg (100k or greater kg) IV every 8 weeks starting 2 weeks after initial load

Refills\*:  None  X6 months  X1 year  Other: \_\_\_\_\_  
 \*(if not indicated order will expire one year from date signed)

*Immunize patients with meningococcal vaccines at least 2 weeks prior to administering the first dose of ULTOMIRIS, unless the risks of delaying ULTOMIRIS therapy outweigh the risk of developing a meningococcal infection. Comply with the most current National Advisory Committee on Immunization (NACI) recommendations for meningococcal vaccination in patients with complement deficiencies.*

Provider Name (Print) \_\_\_\_\_ Physician Signature: \_\_\_\_\_ Date: \_\_\_\_\_

**Fax referral to 866-507-1164 or email to MICreferral@metroinfusioncenter.com**

All information contained in this order form is strictly confidential and will become part of the patient's medical record.