

SOLIRIS (ECULIZUMAB) AND BIOSIMILAR

☐ Soliris (eculizumab)

☐ Bkerv (eculizumab-aeeb)

☐ Dispense As Written (DAW) Will substitute for biosimilar or reference product based on insurance and availability

PATIENT INFORMATION

Patient Name: _____ DOB: _____

Mobile Number: _____ Patient Weight: _____ kg

Allergies: _____

☐ Patient has known difficult venous access. Comments: _____

☐ New Start Therapy | ☐ Continuation of Therapy & Date of last dose (if applicable): _____

DIAGNOSIS (Provider must specify)

☐ Atypical Hemolytic Uremic Syndrome (aHUS), ICD 10: D59.39

☐ Generalized Myasthenia Gravis (gMG), AChR Ab+ (without acute exacerbation), ICD 10: G70.00

☐ Generalized Myasthenia Gravis (gMG), AChR Ab+ (with acute exacerbation), ICD 10: G70.01

☐ Paroxysmal Nocturnal Hemoglobinuria (PNH), ICD 10: D59.5

☐ Neuromyelitis optica spectrum disorder (NMOSD) who are anti-aquaporin-4 (AQP4) antibody positive, ICD 10: G36.0

☐ Other: _____

PROVIDER INFORMATION

Provider Name (print name): _____ Provider NPI: _____

Signature: _____ Date: _____

Contact Name: _____ Phone: _____ Fax: _____

Email Address: _____

Prerequisites to treatment – ensure the following information is complete and attached with referral:

☐ Demographics

☐ Labs/tests supporting diagnosis

☐ Meningococcal vaccination $\geq$ 2 weeks prior to first dose
(Must be current per ACIP with both MenACWY & MenB)

PRE-MEDICATION (Not typically indicated)

☐ Acetaminophen (Tylenol) 500 mg PO

☐ Famotidine 20 mg IV

☐ Methylprednisolone

(Solu-Medrol) 125 mg IVP

☐ Benadryl 25mg PO

☐ Cetirizine (Zyrtec) 10 mg PO

☐ Other: _____ Dose: _____ Route: _____

MEDICATION

MEDICATION	INDICATION	ADULT DOSE	ROUTE
Soliris	PNH	600 mg weekly x 4, then 900 mg 1 week later on week 5, and then 900 mg every 2 weeks thereafter	IV
	gMG, NMOSD, aHUS	900 mg weekly x 4, then 1200 mg 1 week later on week 5, and then 1200 mg every 2 weeks thereafter	

OTHER NOTES

Order valid for 1 year from date of signature unless otherwise specified here: _____