

Patient Information

Patient Name: _____ DOB: _____ Sex: Male Female SSN: _____ Wt (kg/lbs): _____ Ht (cm/in): _____
 Address: _____ Phone: _____ Alternate: _____
 Caregiver Name: _____ Relation to Patient: _____ Phone: _____
 Insurance Plan: _____ Plan ID: _____ BIN #: _____ PCN #: _____ GRP #: _____

Please fax a copy of the front and back of the insurance card(s).

Prescriber + Shipping Information

Prescriber Name: _____ DEA: _____ NPI: _____
 Address: _____
 Phone: _____ Alternate: _____ Fax: _____ Email: _____
 Needs by Date: _____ Ship to: Patient Office Other: _____

Clinical Information (Please fax all pertinent clinical and lab information)

Diagnosis (ICD-10): D59.5 Paroxysmal Nocturnal Hemoglobinuria (PNH) D59.3 Atypical Hemolytic Uremic Syndrome (aHUS) G70.0 Generalized Myasthenia Gravis (gMG)
 Other Code: _____

Prior Therapy <input type="checkbox"/> Yes <input type="checkbox"/> No	Reason for Discontinuation of Therapy	Start Date	End Date

Comorbidities: _____
 Concomitant Medications: _____
 Allergies: NKDA Other: _____

Diagnosis	Dose and Directions	Quantity	Refills
<input type="checkbox"/> Paroxysmal Nocturnal Hemoglobinuria (PNH)	<input type="checkbox"/> Dose Titration: Month 1: Administer 600 mg via IV infusion every 7 days for 4 weeks	<input type="checkbox"/> 4-week supply	0
	<input type="checkbox"/> Maintenance Dosing: Administer 900mg mg via IV INFUSION every 2 weeks starting week 5	<input type="checkbox"/> 4-week supply <input type="checkbox"/> 12-week supply	<input type="checkbox"/> 1 year supply
<input type="checkbox"/> Atypical Hemolytic Uremic Syndrome (aHUS)	<input type="checkbox"/> Dose Titration: Month 1: Administer 900 mg via IV infusion every 7 days for 4 weeks	<input type="checkbox"/> 4-week supply	0
	<input type="checkbox"/> Maintenance Dosing: Administer 1,200 mg via IV infusion every 2 weeks starting week fve	<input type="checkbox"/> 4-week supply <input type="checkbox"/> 12-week supply	<input type="checkbox"/> 1 year supply
<input type="checkbox"/> Generalized Myasthenia Gravis (gMG)	<input type="checkbox"/> Dose Titration: Month 1: Administer 900 mg via IV infusion every 7 days for 4 weeks	<input type="checkbox"/> 4-week supply	0
	<input type="checkbox"/> Maintenance Dosing: Administer 1,200mg via IV infusion every 2 weeks starting week 5	<input type="checkbox"/> 4-week supply <input type="checkbox"/> 12-week supply	<input type="checkbox"/> 1 year supply

Preparation for Administration
 Add eculizumab to an infusion bag and dilute with an equal volume of D5W, sodium chloride 0.9%, sodium chloride 0.45%, or Ringer's injection to a final concentration of 5 mg/mL (eg, 300 mg to a final volume of 60 mL, 600 mg to a final volume of 120 mL, 900 mg to a final volume of 180 mL, or 1,200 mg to a final volume of 240 mL). Gently invert bag to mix thoroughly; do not shake.

Assess vaccination status prior to initiation; patients must receive meningococcal vaccine at least 2 weeks prior to treatment initiation.

Meningococcal Conjugate Vaccine		
<input type="checkbox"/> Menactra®	<input type="checkbox"/> (Up to 55 years) 0.5 mL IM single dose	<input type="checkbox"/> #1
<input type="checkbox"/> Menveo®	<input type="checkbox"/> (Increased risk of infection) 0.5 mL IM as a 2-dose series given at least 8 weeks apart	<input type="checkbox"/> #2
Serogroup B Meningococcal Vaccine		
<input type="checkbox"/> Bexsero®	<input type="checkbox"/> (25 years or younger) 0.5 mL IM as a 2-dose series 1 month apart	<input type="checkbox"/> #2
<input type="checkbox"/> Trumenba®	<input type="checkbox"/> (25 years or younger) 0.5 mL IM as a 2-dose series at 0, and 6 months	<input type="checkbox"/> #2
	<input type="checkbox"/> (25 years or younger) 0.5 mL IM as a 3-dose series at 0, 1 to 2, and 6 months	<input type="checkbox"/> #3

Prescriber's Signature: _____ Date: _____

I authorize AmeriPharma and its representatives to act as an agent to initiate and execute the insurance prior authorization process for this prescription and any future fills of the same prescription for the patient listed above. I understand that I can revoke this designation at any time by providing written notice to AmeriPharma.

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