Lexington 1792 Alysheba Way Suite 205 Lexington, KY 40509

Bowling Green 727 U.S. 31 W Bypass Suite 102 Bowling Green, KY 42101





	\mathbf{IV} (eculizumab-ae	ED) OR				ıte:	
Name:			PATIENT I	NFUKMA	DOB:	SEY.	M □ F □
□NKDA	Allergies:		rnone:		DOB:	Weight lbs/kg:	
	9		DLIVCICIAN	INFORM	ATION		
Physician Name			PHYSICIAN	Practice Nam			
Physician Name: Address:			Office Contact Name: Office Contact #:				
Phone: Fax:			Email (for updates):				
			REFERRA				
□New Referral □Referral Renewal □Medication/Order Cha							
MEDICATION ORDERS □ gMG who are anti-acetylcholine receptor (ArchR) antibody+ ICD 10: gMG NEW START dosing (adult dosing) □ 900 mg weekly for the first 4 weeks, followed by □ 1,200 mg for the fifth dose 1 week later then □ 1,200 mg every 2 weeks x • Refills* □None □x6 months □x1year □Other: *(if not indicated order will expire one year from date signed			□ atypical Hemolytic Uremic Syndrome (aHUS) ICD 10: aHus NEW START dosing (18 yo and older)* □ 900 mg weekly for the first 4 weeks, followed by □ 1,200 mg for the fifth dose 1 week later then □ 1,200 mg every 2 weeks x • Refills* □None □x6 months □x1year □Other: *(if not indicated order will expire one year from date signed PT wt and dosing Body WT Introduction Maintenance				
							□ Paroxysmal Nocturnal Hemoglobinuria (PNH)
			□ WT	30kg to < 40kg	600mg weekly x 2 doses	900mg at week 3 then 900mg every 2 weeks	
PNH BKEMV NEW START dosing (18 yo and older) = 600 mg weekly for the first 4 weeks, followed by				□ WT	20kg to < 30kg	600mg weekly x 2 doses	600mg at week 3 then 600mg every 2 weeks
				□ WT	10kg to < 20kg	600mg weekly x 1 doses	300mg at week 2 then 300mg every 2 weeks
□ 900 mg for the fifth dose 1 week later then □ 900 mg every 2 weeks x • Refills* □None □ x6 months □ x1year □Other: *(if not indicated order will expire one year from date signed			WT	5kg to <10kg	300mg weekly x 1 doses	300mg at week 2 then 300mg every 3 weeks	
			***Complete or update vaccination from meningococcal bacteria (for serogroups A,C,W,Y, and B) at least 2 weeks prior to the first dose of BKEMV, unless the risks of delaying therapy with BKEMV outweigh the risk of developing a serious infection. Comply with the most current Advisory Committee on Immunization Practices (ACIP) recommendations for vaccinations against meningococcal bacteria in patients receiving a complement inhibitor. See Warning and Precautions (5.1) for additional guidance on the management of the risk of serious infections caused by				
REQUIRED DOCUMENTATION CHECKLIST:							
□ This signed order form by the provider □ Patient demographics AND insurance information □ Clinical/ Progress notes supporting primary dx □ Acetylcholine Receptor Antibody Test Results (if Myasthenia Gravis)							
			meningococcal bacteria.				
			WARNINGS AND PRECAUTIONS https://www.accessdata.fda.gov/drugsatfda_docs/label/2024/761333s001lbl.pdf				
□WITH DATE □WITH DATE □IF NOT FUL	of meningococcal vaccines S OF ADMINISTRATION O S OF ADMINISTRATION O LY VACCINATED PROPHYI nt enrolled in the BKEMV RI	f men b & f men ab .actic ai	CWY OR NTIBX RM MUST	BE SENT	enrolled to start	therapy)	

ORDERING PROVIDER

Signature X		Date
Provider	Phone	Provider NPI

□ Is the ordering PROVIDER enrolled in the BKEMV REMS program □YES □NO (If no, must be enrolled to start therapy)