

Oregon Health & Science University Hospital and Clinics Provider's Orders



ADULT AMBULATORY INFUSION ORDER **Eculizumab (SOLIRIS) Infusion**

Page 1 of 4

ACCOUNT NO. MED. REC. NO. NAME BIRTHDATE

Patient Identification

ALL ORDERS MUST BE MARKED IN INK WITH A CHECKMARK (✓) TO BE ACTIVE.

Weight	::	kg Height:cm
Allergi	es:	
Diagno	sis Co	e:
Treatm	ent Sta	t Date: Patient to follow up with provider on date:
This	plan w	Il expire after 365 days at which time a new order will need to be placed
GUIDE	LINES	FOR ORDERING
1.	Send I	ACE SHEET and H&P or most recent chart note.
2.		mab is part of FDA REMS Program
	a.	Providers MUST be enrolled in the SOLIRIS REMS program. MD MUST PROVIDE
		ENROLLMENT ID TO PROCEED:
	b.	Provide patient with both the Patient Safety Brochure and Patient Safety Card. Patient should
		carry the card with them at all times.
	C.	Please see reference links below for enrollment forms and additional help
		i. https://solirisrems.com/Soliris-Prescriber-Enrollment-Form
		iii. https://solirisrems.com/Soliris-Prescriber-Safety-Brochure
		iv. https://solirisrems.com/Soliris-Patient-Safety-Brochure
		v. https://solirisrems.com/Soliris-Patient-Safety-Card
3.	Patien	s must receive the following meningococcal vaccine at least 2 weeks prior to treatment initiation
		Meningococcal serogroups A, C, W, Y vaccine (MenACWY) -Menveo, Menactra, or MenQuad
		These require booster shots every 5 years.
		Date of last vaccination:
	b.	Meningococcal serogroup B vaccine -Bexsero or Trumenba. These require booster shots 1 ye
		after primary series and every 2 to 3 years thereafter.
	_	Date of last vaccination:
		entation for vaccines must be sent with the order.
		s not vaccinated should be on prophylaxis antibiotics until vaccines are up to date. Patients wh een vaccinated less than 2 weeks prior to start of infusion should be on 2 weeks of antibacteria
	nave L	seri vaccinated less trian 2 weeks prior to start or influsion should be on 2 weeks or antibacteria

prophylaxis.

Prescriber must update the status of the patient's meningococcal vaccination, indication, and antibacterial drug prophylaxis into the ultsorems.com online portal.

Has this been done? (Yes) (No) - Circle One

If no, you must document the one-time status into the online portal before you may treat the patient.

- 4. Treatment should be administered at the recommended time interval although administration may vary by ±2 days.
- 5. Monitoring during therapy: monitor platelet count, serum LDH levels, and serum creatinine levels during therapy. Monitor for signs and symptoms of infection, in particular meningococcal infections.
- 6. Monitoring after discontinuation:
 - a. Atypical hemolytic uremic syndrome (aHUS) patients who discontinue treatment should be monitored closely for at least 12 weeks for signs and symptoms of thrombotic microangiopathy (TMA) complications.



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Page 2 of 4

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- b. Paroxysmal nocturnal hemoglobinuria (PNH) patients who discontinue treatment should be monitored for at least 8 weeks for signs and symptoms of hemolysis
- 7. Consider penicillin prophylaxis for the duration of eculizumab therapy to potentially reduce the risk of meningococcal disease.

PR		CREENING: (Must be available prior to initiation of therapy):
	•	Meningococcal serogroups A, C, W, Y vaccine (MenACWY) -MenQuadfi, Menactra, or Menveo given on (dates)
	•	Meningococcal serogroup B vaccine -Bexsero or Trumenba given on (dates)
LAI	3S:	
		CBC with differential, Routine, ONCE, every (visit)(days)(weeks)(months) – Circle One CMP, Routine, ONCE, every (visit)(days)(weeks)(months) – Circle One LDH TOTAL, Routine, ONCE every (visit)(days)(weeks)(months) – Circle One
		Labs already drawn. Date: (visit)(days)(weeks)(months) = Circle Orie
NU	RSI	NG ORDERS:
		Vital signs at baseline, post-infusion, and prior to discharge.
		Monitor for 1 hour after infusion complete for signs or symptoms of infusion reaction. May discontinue observation if stable and tolerating infusions.
	3.	Hold treatment and notify provider if patient is not up to date on meningococcal vaccination every 5 years for MenACWY (Menveo, Menactra, or MenQuadfi) or 1 year after primary series and every 2 to 3 years thereafter for MenB (either Bexsero or Trumenba).
	4.	Follow facility policies and/or protocols for vascular access maintenance with appropriate flush solution, declotting (alteplase), and/or dressing changes.
ME	DIC	ATIONS:
		Atypical hemolytic uremic syndrome (aHUS), Generalized myasthenia gravis, refractory, or
		Neuromyelitis optica spectrum disorder

N

	Initial doses : eculizumab (SOLIRIS) 900 mg in sodium chloride 0.9% 90 mL, intravenous, ONCE Every week x 4 doses
	Maintenance doses : eculizumab (SOLIRIS) 1200 mg in sodium chloride 0.9% 120 mL, intravenous, ONCE Every 2 weeks x doses, begin on week 5
Inf	use over 35 minutes. Infusion may be slowed or stopped due to adverse reactions but should be

finished within 2 hours Provide patient with Soliris Patient Safety Card to keep at all times



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Page 3 of 4

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<u>P</u> a	<u>aro)</u>	oxysmal nocturnal hemoglobinuria (PNH)	
		☐ Initial doses: eculizumab (SOLIRIS) 600 mg in NaCl 0.9% 60 mL, intravenous, ONCE Every week x 4 doses	
		☐ Maintenance doses : eculizumab (SOLIRIS) 900 mg in NaCl 0.9% 90 mL, intravenous, ON Every 2 weeks x doses, begin on week 5	CE
	fini	nfuse over 35 minutes. Infusion may be slowed or stopped due to adverse reactions but should inished within 2 hours Provide patient with Soliris Patient Safety Card to keep at all times	d be
 HYPERSENSITIVITY MEDICATIONS: NURSING COMMUNICATION – If hypersensitivity or infusion reactions develop, temporarily infusion and notify provider immediately. Administer emergency medications per the Treatmed Algorithm for Acute Infusion Reaction (OHSU HC-PAT-133-GUD, HMC C-132). Refer to algo symptom monitoring and continuously assess as grade of severity may progress. diphenhydrAMINE (BENADRYL) injection, 25-50 mg, intravenous, AS NEEDED x 1 dose for hypersensitivity or infusion reaction EPINEPHrine HCI (ADRENALIN) injection, 0.3 mg, intramuscular, AS NEEDED x 1 dose for hypersensitivity or infusion reaction hydrocortisone sodium succinate (SOLU-CORTEF) injection, 100 mg, intravenous, AS NEEDED dose for hypersensitivity or infusion reaction famotidine (PEPCID) injection, 20 mg, intravenous, AS NEEDED x 1 dose for hypersensitivity infusion reaction 			
I am re I hold a that co	espo an a orres	ing below, I represent the following: ponsible for the care of the patient (who is identified at the top of this form); n active, unrestricted license to practice medicine in: □ Oregon □ (chec responds with state where you provide care to patient and where you are currently licensed. Spatient Oregon);	k box secify
My ph PRESO medica	ysic CRI ation	(MUST BE COMPLETED TO BE A VALID RIPTION); and I am acting within my scope of practice and authorized by law to order Infusion described above for the patient identified on this form.	of the
Provi	der	er signature: Date/Time:	
Print	ed I	d Name: Phone: Fax:	



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Page 4 of 4

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Please check the appropriate box for the patient's preferred clinic location:

☐ Hillsboro Medical Center

Infusion Services 364 SE 8th Ave, Medical Plaza Suite 108B Hillsboro, OR 97123

Phone number: (503) 681-4124 Fax number: (503) 681-4120

☐ Mid-Columbia Medical Center

Celilo Cancer Center 1800 E 19th St The Dalles, OR 97058

Phone number: (541) 296-7585 Fax number: (541) 296-7610 ☐ Adventist Health Portland

Infusion Services 10123 SE Market St Portland, OR 97216

Phone number: (503) 261-6631 Fax number: (503) 261-6756