

Empaveli Clinical Profile Fax Form



PANTHERx Rare Pharmacy
121 Bayer Road, Building 5
Pittsburgh, PA 15205
Phone: 866-258-1895 Fax: 866-609-1760

Physician:
Fax Number:

Date/Time:

Patient Name: DOB:

Please use this form to complete the Initial Referral process for EMPAVELI™. Please fax this completed form and any relevant clinical documentation (if applicable) attached to 1-866-609-1760.

If you have any questions, please call 1-866-258-1895 and select Option 2 then Option 3 to speak with a pharmacist.

Vaccine Brand	ACIP Recommendation for Patients with Complement Deficiency	Dose in Series (Administration Date)
Meningococcal Conjugate Vaccine (MenACWY) Vaccination History		
<input type="checkbox"/> Menactra® (MenACWY-D) <input type="checkbox"/> Menveo® (MenACWY-CRM) <input type="checkbox"/> MenQuadfi® (MenACWY-TT)	Administer a 2-dose series of MenACWY (Menactra, Menveo, or MenQuadfi) at least 8 weeks apart and revaccinate every 5 years if risk remains.	<input type="checkbox"/> Dose #1 (Date: _____) <input type="checkbox"/> Dose #2 (Date: _____) <input type="checkbox"/> Boosters (if applicable) (Most Recent Dose Date: _____)
Serogroup B Meningococcal Vaccination History		
<input type="checkbox"/> Bexsero® (MenB-4C) <input type="checkbox"/> Trumenba® (MenB-FHbp)	Administer 2-dose primary series MenB-4C (Bexsero) at least 1 month apart or 3-dose primary series MenB-FHbp (Trumenba) at 0, 1-2, 6 months (if dose 2 was administered at least 6 months after dose 1, dose 3 not needed. MenB-4C and MenB-FHbp are not interchangeable (use same product for all doses in series). 1 dose MenB booster 1 year after primary series and revaccinate every 2-3 years if risk remains.	<input type="checkbox"/> Dose #1 (Date: _____) <input type="checkbox"/> Dose #2 (Date: _____) <input type="checkbox"/> Dose #3 (if applicable) (Date: _____) <input type="checkbox"/> Boosters (if applicable) (Most Recent Dose Date: _____)
Pneumococcal PCV13 Vaccination History		
<input type="checkbox"/> Prevnar 13® (PCV13)	1 dose PCV13 followed by 1 dose PPSV23 at least 8 weeks later, then another dose PPSV23 at least 5 years after previous PPSV23; at age 65 years or older, administer 1 dose PPSV23 at least 5 years after most PPSV23 (note: only 1 dose PPSV23 recommended at age 65 years or older.	<input type="checkbox"/> Dose #1 (Date: _____)
Pneumococcal PPSV23 Vaccination History		

<input type="checkbox"/> Pneumovax® (PPSV23)	1 dose PCV13 followed by 1 dose PPSV23 at least 8 weeks later, then another dose PPSV23 at least 5 years after previous PPSV23; at age 65 years or older, administer 1 dose PPSV23 at least 5 years after most PPSV23 (note: only 1 dose PPSV23 recommended at age 65 years or older.	<input type="checkbox"/> Dose #1 (Date: _____) <input type="checkbox"/> Dose #2 (if applicable) (Date: _____) <input type="checkbox"/> Dose #3 (if applicable) (Date: _____)
Haemophilus influenza type B (Hib) Vaccination History		
<input type="checkbox"/> ActHIB® <input type="checkbox"/> Hiberix® <input type="checkbox"/> PedvaxHIB®	This is a recommended vaccination for adults who meet the age requirement and lack documentation of vaccination or lack evidence of past infection. If patients lack documentation of typical childhood scheduled/catch up vaccination, 1 dose of Hib (ActHIB, Hiberix, or PedvaxHIB)	<input type="checkbox"/> Dose #1 (Date: _____) <input type="checkbox"/> Dose #2 (if applicable) (Date: _____) <input type="checkbox"/> Dose #3 (if applicable) (Date: _____) <input type="checkbox"/> Dose #4 (if applicable) (Date: _____)
1. PNH Diagnosis Date: Date when patient was first diagnosed with PNH (month/day/year) _____		
1. Most Recent/Current Therapy: Current treatment prior to product switch to EMPAVELI™ <input type="checkbox"/> Eculizumab (Soliris®) <input type="checkbox"/> Raculizumab-cwvz (Ultomiris®) <input type="checkbox"/> pegcetacoplan (EMPAVELI™) (clinical trial) <input type="checkbox"/> No prior complement inhibitor therapy prior to planned EMPAVELI™ start <input type="checkbox"/> Other (please specify) _____		
2. Prior Therapy Dose and Frequency: Current treatment dose and frequency prior to product switch to Empaveli <div style="display: flex; justify-content: space-between;"> <div style="width: 45%;"> <u>Dose:</u> <input type="checkbox"/> 900 mg (Soliris®) <input type="checkbox"/> 3,000 mg (40kg to < 60kg Ultomiris®) <input type="checkbox"/> 3,300 mg (60kg to < 100kg Ultomiris®) <input type="checkbox"/> 3,600 mg (≥ 100kg Ultomiris®) <input type="checkbox"/> Other (please specify) _____ mg </div> <div style="width: 45%;"> <u>Frequency:</u> <input type="checkbox"/> Once every 2 weeks <input type="checkbox"/> Once every 8 weeks <input type="checkbox"/> Other (please specify) _____ </div> </div>		
3. Prior Therapy Start: Current therapy start date prior to product switch to EMPAVELI™ _____		
4. Years on Complement Inhibitor Therapy: Number of years on complement inhibitor therapy _____		
5. Current Therapy Most Recent Dose: Date of most recent dose for current therapy _____		
6. Anticipated Start Date of EMPAVELI™: What date do you anticipate the patient will start therapy with EMPAVELI™ (month/day/year) _____		
7. Anticipated Last Dose Date of Current Therapy: Anticipated last scheduled dose of current complement inhibitor therapy, based upon your current targeted start date of EMPAVELI™ (month/day/year) _____		
8. Hemoglobin: Most recent hemoglobin value prior to starting EMPAVELI™ _____ (g/dL)		
9. Bilirubin: Most recent bilirubin value prior to starting EMPAVELI™ _____ (μmol/L)		
10. Reticulocyte Count: Most recent reticulocyte count prior to starting EMPAVELI™ _____ (10x9 cells/L)		
11. LDH level: Most recent lactate dehydrogenase value prior to starting EMPAVELI™ _____ (unit/L)		
12. Transfusion History: Number of transfusions within 6 months prior to starting EMPAVELI™ _____		
13. Breakthrough Hemolysis History: Number of breakthrough hemolysis events within 6 months prior to starting EMPAVELI™ _____		
14. Thrombosis History: Number of thrombotic events within 6 months prior to starting EMPAVELI™ _____		

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