

COMPLEMENT INHIBITORS PRIOR AUTHORIZATION FORM (form effective 07/01/2022)

Prior authorization guidelines for **Complement Inhibitors** and **Quantity Limits** are available on the DHS Pharmacy Services website at <https://www.dhs.pa.gov/providers/Pharmacy-Services/Pages/default.aspx>.

<input type="checkbox"/> New request <input type="checkbox"/> Renewal request		# of pages: _____	Prescriber name:	
Name of office contact:			Specialty:	
Contact's phone number:			NPI:	State license #:
LTC facility contact/phone:			Street:	
Beneficiary name:			City/state/zip:	
Beneficiary ID#:	DOB:	Phone:	Fax:	

CLINICAL INFORMATION

Drug requested – LOADING dose:	Drug requested – MAINTENANCE dose:
<input type="checkbox"/> Enjaymo** – Strength & package size: _____ # of vials per dose: _____ # of doses/refills requested: _____	<input type="checkbox"/> Empaveli – Strength & package size: _____ # of vials per dose: _____ # of doses/refills requested: _____
<input type="checkbox"/> Soliris** – Strength & package size: _____ # of vials per dose: _____ # of doses/refills requested: _____	<input type="checkbox"/> Enjaymo** – Strength & package size: _____ # of vials per dose: _____ # of doses/refills requested: _____
<input type="checkbox"/> Ultomiris** – Strength & package size: _____ # of vials per dose: _____ # of doses/refills requested: _____	<input type="checkbox"/> Soliris** – Strength & package size: _____ # of vials per dose: _____ # of doses/refills requested: _____
<input type="checkbox"/> other: _____ _____ _____	<input type="checkbox"/> Tavneos capsule – Strength: _____ Quantity: _____ Refills requested: _____
<p>**Indicates drug is included in the Specialty Pharmacy Drug Program. Further information included below.</p>	<input type="checkbox"/> Ultomiris** – Strength & package size: _____ # of vials per dose: _____ Refills requested: _____
<input type="checkbox"/> other: _____	<input type="checkbox"/> other: _____
LOADING dose/directions:	MAINTENANCE dose/directions:

Diagnosis (<i>submit documentation</i>):	Dx code (<i>required</i>):	Weight (kg):
**SPECIALTY PHARMACY DRUG PROGRAM: <u>Enjaymo</u> , <u>Soliris</u> , and <u>Ultomiris</u> are included in the DHS Specialty Pharmacy Drug Program and is available from DHS's specialty pharmacy. Refer to https://www.dhs.pa.gov/providers/Pharmacy-Services/Pages/Specialty-Pharmacy-Program.aspx for more information about the Specialty Pharmacy Drug Program.		DHS specialty pharmacy: Chartwell Pennsylvania, LP Oakdale, PA Phone: 833-710-0211 Fax: 412-920-1869 www.chartwellpa.com
Is the requested medication being prescribed by or in consultation with a specialist (e.g., hematologist/oncologist, neurologist, nephrologist, etc.)?	<input type="checkbox"/> Yes <input type="checkbox"/> No	<i>Submit documentation of consultation if applicable.</i>

INITIAL requests

Check all of the following that apply to the beneficiary and this request and **SUBMIT DOCUMENTATION** for each item.

- The beneficiary is up to date on the following vaccines as recommended in the FDA-approved package labeling:
 - meningococcal conjugate (MenACWY; Menactra, Menveo)
 - meningococcal serogroup B (MenB; Bexsero, Trumenba)
 - pneumococcal conjugate (PCV13; Prevnar 13)
 - pneumococcal polysaccharide (PPSV23; Pneumovax 23)
 - Haemophilus influenza (Hib)
- For the treatment of generalized myasthenia gravis:
 - Has a positive serologic test for anti-AChR antibodies
 - Has a Myasthenia Gravis Foundation of America (MGFA) Clinical Classification of class II, III, or IV at initiation of therapy
 - Has a myasthenia gravis-specific activities of daily living scale (MG-ADL) total score ≥ 6 at initiation of therapy
 - Failed treatment over 6 months or more with 2 or more immunosuppressive therapies either in combination or as monotherapy
 - Has a contraindication or intolerance to immunosuppressive therapies
 - Failed treatment with IVIG
 - Has a contraindication or intolerance to IVIG
 - Failed treatment with plasma exchange
 - Has a contraindication or intolerance to plasma exchange
- For the treatment of neuromyelitis optica spectrum disorder:
 - Failed treatment with rituximab
 - Has a contraindication or intolerance to rituximab
- For Empaveli (pegcetacoplan):
 - Has results of a recent lactate dehydrogenase level
 - Prescribed dose is appropriate based on recent lactate dehydrogenase level
- For Tavneos (avacopan):
 - Prescribed dose is appropriate based on concomitant strong CYP3A4 inhibitors (e.g., protease inhibitors, azole antifungals, nefazodone)
 - Not taking concomitant strong CYP3A4 inhibitors (*submit complete medication list*)

RENEWAL requests

Check all of the following that apply to the beneficiary and this request and **SUBMIT DOCUMENTATION** for each item.

- Tolerated the requested medication (no significant adverse effects)
- Experienced a positive clinical response to the requested medication
- For Empaveli (pegcetacoplan):
 - Has results of a recent lactate dehydrogenase level
 - Prescribed dose is appropriate based on recent lactate dehydrogenase level

- For Tavneos (avacopan):
- Prescribed dose is appropriate based on concomitant strong CYP3A4 inhibitors (e.g., protease inhibitors, azole antifungals, nefazodone)
 - Not taking concomitant strong CYP3A4 inhibitors (*submit complete medication list*)

PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION TO DHS – PHARMACY DIVISION

Prescriber Signature:

Date:

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