Field Name	Field Description
Prior Authorization	Veopoz
Group Description	
Drugs	Veopoz (pozelimab-bbfg)
Covered Uses	Medically accepted indications are defined using the following sources: the Food and Drug Administration (FDA), Micromedex, American Hospital Formulary Service (AHFS), United States Pharmacopeia Drug Information for the Healthcare Professional (USP DI), the Drug Package Insert (PPI), or disease state specific standard of care guidelines.
Exclusion Criteria	 Patients with unresolved Neisseria meningitidis infection Concurrent use of another complement inhibitor (i.e. Soliris)
Required Medical Information	See "Other Criteria"
Age Restrictions	According to package insert
Prescriber Restrictions	Prescribed by or in consultation with a physician with experience in managing complement related disorders (i.e., gastroenterologist, immunologist, cardiologist, etc.)
Coverage Duration	If all of the criteria are met, the initial request will be approved for 6 months. For continuation of therapy, the request will be approved for 12 months.
Other Criteria	Initial Authorization:
	 Medication is prescribed at an FDA approved dose Diagnosis of CD55-deficient protein-losing enteropathy (PLE), also known as CHAPLE disease Documentation of hypoalbuminemia (serum albumin <3.5 g/dL) Documentation of patient weight Re-Authorization:
Revision/Review Date: 11/2024	 Documentation or provider attestation of positive clinical response (i.e. symptom improvement, normalization of labs such as serum albumin (3.5-5.5 g/dL) and IgG concentrations, reduced hospitalizations and severe adverse events, increased quality of life, etc.) Documentation of patient weight Medication is prescribed at an FDA approved dose
	If all of the above criteria are not met, the request is referred to a Medical Director/Clinical Reviewer for medical necessity review.