

**Veopoz**  
**Effective 03/01/2024**

<b>Plan</b>	<input type="checkbox"/> MassHealth UPPL <input checked="" type="checkbox"/> Commercial/Exchange	<b>Program Type</b>	<input checked="" type="checkbox"/> Prior Authorization <input type="checkbox"/> Quantity Limit <input type="checkbox"/> Step Therapy
<b>Benefit</b>	<input type="checkbox"/> Pharmacy Benefit <input checked="" type="checkbox"/> Medical Benefit		
<b>Specialty Limitations</b>	N/A		
<b>Contact Information</b>	<b>Specialty Medications</b>		
	All Plans	Phone: 877-519-1908	Fax: 855-540-3693
<b>Exceptions</b>	<b>Non-Specialty Medications</b>		
	All Plans	Phone: 800-711-4555	Fax: 844-403-1029

**Overview**

Veopoz is indicated for the treatment of adult and pediatric patients 1 year of age and older with CD55-deficient protein-losing enteropathy (PLE), also known as CHAPLE disease.

**Coverage Guidelines**

Authorization may be granted for members new to the plan who are currently receiving treatment with the requested medication, excluding when the product is obtained as samples or via manufacturer’s patient assistance programs.

**OR**

Authorization may be granted when the following criteria is met:

1. The member has a diagnosis of CD55-deficient protein-losing enteropathy (PLE)
2. Medical records documenting ALL of the following:
  - a. The member has confirmed biallelic CD55 loss-of-function mutation detected by genotype analysis.
  - b. The member has hypoalbuminemia (serum albumin concentration of  $\leq 3.2$  g/dL)
  - c. The member has ONE of the more of the following signs and symptoms of CD-55 PLE within the past 6 months:
    - i. Abdominal pain
    - ii. Diarrhea
    - iii. Peripheral edema
    - iv. Facial edema

**Continuation of Therapy**

Authorization may be granted for continued treatment in members when there is no evidence of unacceptable toxicity or disease progression AND member demonstrates a positive response to therapy (e.g., normalization of serum albumin, improvement in signs and symptoms of disease and/or decrease in number of hospitalizations and infections)

**Limitations**

1. Initial approvals will be granted for 6 months.
2. Reauthorizations will be granted for 12 months.

**References**

1. Veopoz [package insert]. Tarrytown, NY: Regeneron Pharmaceuticals, Inc.; August 2023.

**Review History**

2/14/2023: Created and Reviewed at Feb P&T, Effective 3/1/2024

