

Fabhalta (iptacopan)
Effective 10/1/2024

Plan	<input type="checkbox"/> MassHealth UPPL <input checked="" type="checkbox"/> Commercial/Exchange	Program Type	<input checked="" type="checkbox"/> Prior Authorization <input type="checkbox"/> Quantity Limit <input type="checkbox"/> Step Therapy
Benefit	<input checked="" type="checkbox"/> Pharmacy Benefit <input type="checkbox"/> Medical Benefit		
Specialty Limitations	This medication has been designated specialty and must be filled at a contracted specialty pharmacy.		
Contact Information	Medical and Specialty Medications		
	All Plans	Phone: 877-519-1908	Fax: 855-540-3693
Exceptions	Non-Specialty Medications		
	All Plans	Phone: 800-711-4555	Fax: 844-403-1029
Exceptions	N/A		

Overview

Fabhalta (iptacopan) is a complement factor B inhibitor indicated for the treatment of adults with paroxysmal nocturnal hemoglobinuria.

Coverage Guidelines

Authorization may be granted for members new to the plan within the previous 90 days 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 if the member meets all of the following criteria and documentation has been submitted:

- The member has a diagnosis of paroxysmal nocturnal hemoglobinuria confirmed by flow cytometry

Continuation of Therapy

Reauthorization will be granted when the following criteria are met:

- Prescriber submits documentation of a positive response to therapy (e.g., normalization of lactate dehydrogenase [LDH] levels, hemoglobin stabilization, decreased number of red blood cell transfusions)

Limitations

- Initial and reauthorization approvals will be granted for 12 months.

References

- Fabhalta (iptacopan) capsules [prescribing information]. East Hanover, NJ: Novartis Pharmaceuticals Corporation; March 2024.

Review History

08/14/2024 – Reviewed at August P&T. Effective 10/1/2024.

