

Hematopoietics/Chronic ITP Prior Authorization of Benefits Form

CONTAINS CONFIDENTIAL PATIENT INFORMATION

Complete form in its entirety and fax to: Prior Authorization of Benefits Center at 1-844-512-9004. Provider Help Desk: 1-800-454-3730

3. Medication	4. Strength	5. Directions	6. Quantity per 30 days
			Specify:
7. Diagnosis:			

8. Approval criteria: (Check all boxes that apply. Note: Any areas not filled out are considered not applicable to your patient and may affect the outcome of this request.)

Prior authorization is required for hematopoietics/chronic ITP agents. Request must adhere to all FDA-approved labeling. Payment for a nonpreferred hematopoietic/chronic ITP agent will be considered following documentation of a recent trial and therapy failure with a preferred hematopoietic/chronic ITP agent, when applicable, unless such a trial would be medically contraindicated. Payment will be considered under the following conditions:

Preferred	<u>Nonpreferred</u>
🗆 Promacta	□ Doptelet
	Mulpleta
	🗆 Nplate
	🗆 Promacta powder
	nia with chronic immune thrombocytopenia (ITP) (Doptelet, Promacta, Nplate, Tavalisse) an insufficient response to a corticosteroid, immunoglobulin, or splenectomy.
Documentation of a	
Trial drug name:	
Trial start date:	Trial end date:

Tria Failure reason: Has the patient undergone splenectomy? \Box No \Box Yes

🗆 Severe aplastic anemia (Promacta)			
1. Patient has documentation of an insufficient response or intolerance to at least one prior immunosuppressive therapy;			
and			
2. Patient has a platelet count \leq 30 x 10 ⁹ /L.			
3. If criteria for coverage are met, initial authorization will be given for 16 weeks.			
Documentation of hematologic response after 16 weeks of therapy will be required for further consideration.			
Trial drug name:			
Trial start date:Trial end date:			
Failure reason:			
Platelet count: Lab date:			
Renewal requests:			
Has patient has a hematologic response after 16 weeks of Promacta therapy? 🗆 Yes (attach labs) 🛛 No			
□ Thrombocytopenia with chronic liver disease in patients scheduled to undergo a procedure (Doptelet, Mulpleta)			
Documentation of the following:			
1. Pre-treatment platelet count; and			
2. Scheduled dosing prior to procedure; and			
3. Therapy completion prior to scheduled procedure; and			
4. Platelet count will be obtained before procedure.			
Platelet count: Lab date:			
Date of scheduled procedure:			
Date for start of drug treatment:			
After the last dose, a platelet count will be obtained prior to undergoing the procedure: \Box Yes \Box No			
Other diagnosis:			
Reason for use of Non-Preferred drug requiring prior approval:			
Other medical conditions to consider:			
Attach lab results and other documentation as necessary.			

9. Physician signature

Prescriber or authorized signature

Date

Prior Authorization of Benefits is not the practice of medicine or the substitute for the independent medical judgment of a treating physician. Only a treating physician can determine what medications are appropriate for a patient. Please refer to the applicable plan for the detailed information regarding benefits, conditions, limitations, and exclusions. The submitting provider certifies that the information provided is true, accurate, and complete and the requested services are medically indicated and necessary to the health of the patient.

Note: Payment is subject to member eligibility. Authorization does not guarantee payment.