

HEALTH PARTNERS PLANS PRIOR AUTHORIZATION REQUEST FORM

Thrombopoietics

Phone: 215-991-4300 Fax back to: 866-240-3712

Health Partners Plans manages the pharmacy drug benefit for your patient. Certain requests for coverage require review with the prescribing physician. Please answer the following questions and fax this form to the number listed above.

PLEASE NOTE: Any information (patient, prescriber, drug, labs) left blank, illegible, or not attached WILL DELAY the review process.

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Patient Name:		Prescriber Name:		
HPP Member Number:		Fax:	Phone:	
Date of Birth:		Office Contact:		
Patient Primary Phone:		NPI:	PA PROMISe ID:	
Address:		Address:		
City, State ZIP:		City, State ZIP:		
Line of Business: ☐ Medicaid ☐ CHIP		Specialty Pharmacy (if applicable):		
Drug Name:		Strength:		
Quantity:		Refills:		
Directions:				
Diagnosis Code:	Diagnosis:			
-		onths but may be less dependir	 na on the drua	
This emaximum approx	var timo io 12 mi	shine but may be less depending	g on the drug.	
Please attach any pertinent medical histor	ry including lab	os and information for this me	mber that may support approval.	
Please	answer the fol	llowing questions and sign.		
Q1. Is this a request for continuation of therapy with the requested product? [If NO, skip to 13.]				
Yes	□ No			
Q2. Is the requested product prescribed for thrombocytopenia in a patient scheduled to undergo a procedure that was previously approved?				
☐ Yes ☐ No				
Q3. Is this request for Doptelet (avatrombopag) or Mulpleta (lusutrombopag [If YES, skip to 14.]				
☐ Yes ☐ No				
Q4. Is the requested product prescribed by or in consultation with an appropriate specialist (i.e., hematologist/oncologist, gastroenterologist, hepatologist, etc.)?				
☐ Yes ☐ No				
Q5. Is the prescribed dose and duration of therapy consistent with Food and Drug Administration (FDA)-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature?				
☐ Yes ☐ No				
Q6. Has the patient had a documented increase in platelet count sufficient to avoid bleeding that requires medical attention?				
☐ Yes	☐ Yes ☐ No			

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Patient Name:	Prescriber Name:		
Q7. Is this request for the treatment of severe aplastic anemia?			
☐ Yes	□ No		
Q8. Is there documentation of a positive clinical response to the requested product?			
☐ Yes	□No		
Q9. Is there documentation of repeat lab results and monitoring as recommended in the Food and Drug Administration (FDA)-approved package labeling?			
☐ Yes	□No		
Q10. Is this request for Tavalisse (fostamatinib)?			
☐ Yes	□ No		
Q11. Does the patient have diarrhea that is greater than or equal to grade 3?			
☐ Yes	□ No		
Q12. Is there a documented plan to manage the diarrhea that is consistent with Food and Drug Administration (FDA)-approved package labeling?			
☐ Yes	□ No		
Q13. Is this request for Doptelet (avatrombopag), Mulpleta (lusutrombopag), or Tavalisse (fostamatinib)?			
☐ Yes	□ No		
Q14. Does the patient have documented therapeutic failure, contraindication, or intolerance to the preferred thrombopoietics approved for the patient's indication?			
☐ Yes	□ No		
Q15. Is this request for Nplate (romiplostim) or Promacta (eltrombopag)?			
☐ Yes	□ No		
Q16. Is the requested product prescribed by or in consultation with an appropriate specialist (i.e., hematologist/oncologist, gastroenterologist, hepatologist, etc.)?			
☐ Yes	□ No		
Q17. Is the patient being treated for a diagnosis that is indicated in the Food and Drug Administration (FDA)-approved package labeling OR a medically accepted indication?			
☐ Yes	□ No		
Q18. Is the prescribed dose and duration of therapy consistent with Food and Drug Administration (FDA)-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature?			

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Patient Name:	Prescriber Name:	
☐ Yes	□No	
Q19. Is there documentation of baseline lab results and n Administration (FDA)-approved package labeling?	nonitoring as recommended in the Food and Drug	
☐ Yes	□ No	
Q20. Is the requested drug prescribed for the treatment o	f thrombocytopenia prior to a procedure?	
☐ Yes	□ No	
Q21. Does the patient have a documented pretreatment p	platelet count of less than 50,000 cells per microliter?	
☐ Yes	□ No	
Q22. Will the patient begin treatment with the requested proof and Drug Administration (FDA)-approved package	product prior to the scheduled procedure in accordance with abeling?	
☐ Yes	□ No	
Q23. Does the patient have a documented pretreatment p	platelet count of less than 30,000 cells per microliter?	
☐ Yes	□ No	
Q24. Is the requested product prescribed for the treatmer	nt of immune thrombocytopenia (ITP)?	
☐ Yes	□ No	
Q25. Is this request for Promacta (eltrombopag) prescribe	ed for the treatment of refractory severe aplastic anemia?	
☐ Yes	□ No	
Q26. Is this request for Promacta (eltrombopag) prescribe	ed for the primary treatment of aplastic anemia?	
☐ Yes	□ No	
Q27. Additional Information:		
Prescriber Signature	Date	

Updated for 2020