



Commonwealth of Massachusetts
MassHealth Drug Utilization Review Program
 P.O. Box 2586, Worcester, MA 01613-2586
Fax: 1-877-208-7428 **Phone:** 1-800-745-7318

Immune Thrombocytopenia Agents Prior Authorization Request

MassHealth reviews requests for prior authorization (PA) on the basis of medical necessity only. If MassHealth approves the request, payment is still subject to all general conditions of MassHealth, including current member eligibility, other insurance, and program restrictions. MassHealth will notify the requesting provider and member of its decision. Keep a copy of this form for your records. If faxing this form, please use black ink.

Additional information about these agents, including PA requirements and preferred products, can be found within the MassHealth Drug List at www.mass.gov/druglist.

Member information

Last name _____ First name _____ MI _____
 MassHealth member ID # _____ Date of birth _____
 Gender (Check one.) F M Member's place of residence home nursing facility

Medication information

Medication requested

- | | |
|---|---|
| <input type="checkbox"/> Doptelet (avatrombopag) | <input type="checkbox"/> Promacta (eltrombopag) |
| <input type="checkbox"/> Mulpleta (lusutrombopag) | <input type="checkbox"/> Tavalisse (fostamatinib) |
| <input type="checkbox"/> Nplate (romiplostim) | |

Dose and frequency _____ **Duration of therapy** _____

Indication (Check all that apply.)

- | | |
|---|--|
| <input type="checkbox"/> Chronic, relapsed, or refractory immune thrombocytopenia (ITP) | <input type="checkbox"/> Thrombocytopenia due to chronic liver disease (CLD) |
| <input type="checkbox"/> Severe aplastic anemia | <input type="checkbox"/> Thrombocytopenia in the setting of hepatitis C |

Section I. Please complete for Doptelet and Mulpleta requests.

- Is a procedure planned?
 Yes. Please provide anticipated date of procedure. _____ No
- Please provide date and results of most recent platelet count (including laboratory reference ranges).

- For Doptelet requests, has the member had a trial with Mulpleta?
 Yes. Please list the dates/duration of use and outcomes below.
 Dates/duration of use _____ Adverse reaction Inadequate response Other
 Briefly describe details of adverse reaction, inadequate response, or other.

 No. Please explain why not. _____

Section II. Please complete for Nplate, Promacta, and Tavalisse requests for chronic, relapsed or refractory ITP.

1. Please provide date and results of most recent platelet count (including laboratory reference ranges). For platelet count > 30,000 cells/mcL, describe medical necessity for platelet elevation.

2. Has the member had a trial with a corticosteroid?
 Yes. Please list the drug name, dates/duration of use, and outcomes below.
Drug name _____ Dates/duration of use _____
Did the member experience any of the following? Adverse reaction Inadequate response Other
Briefly describe details of adverse reaction, inadequate response, or other.

- No. Please explain why not. _____
3. Has the member had a trial with immunoglobulin therapy?
 Yes. Please list the drug name, dates/duration of use, and outcomes below.
Drug name _____ Dates/duration of use _____
Did the member experience any of the following? Adverse reaction Inadequate response Other
Briefly describe details of adverse reaction, inadequate response, or other.

- No. Please explain why not. _____
4. Has the member had a splenectomy? Yes No
5. For Nplate and Tavalisse requests, has the member had a trial with Promacta?
 Yes. Please list the dates/duration of use and outcomes below.
Dates/duration of use _____ Adverse reaction Inadequate response Other
Briefly describe details of adverse reaction, inadequate response, or other.

- No. Please explain why not. _____

Section III. Please complete for Promacta requests for thrombocytopenia in the setting of hepatitis C.

1. Please provide date and results of most recent platelet count (including laboratory reference ranges).

2. Is the member currently on interferon therapy? Yes. Please provide start date. _____ No
3. For members not currently on interferon therapy, does the treatment plan include initiation of therapy with interferon? Yes No

Section IV. Please complete for Promacta requests for severe aplastic anemia.

1. Please provide date and results of most recent platelet count (including laboratory reference ranges).

2. Has the member had a trial with anti-thymocyte globulin (ATG)?
 Yes. Please list the dates/duration of use and outcomes below.
Dates/duration of use _____ Adverse reaction Inadequate response Other
Briefly describe details of adverse reaction, inadequate response, or other.

- No. Please explain why not. _____
3. Has the member had a trial with cyclosporine?
 Yes. Please list the dates/duration of use and outcomes below.
Dates/duration of use _____ Adverse reaction Inadequate response Other

Briefly describe details of adverse reaction, inadequate response, or other.

No. Please explain why not. _____

4. For use of Promacta in combination with ATG and cyclosporine, please provide clinical rationale.

Section V. Please complete for all requests for non-preferred drug products if one or more preferred drug products have been designated for this class of drugs.

If one or more preferred drug products have been designated for this class of drugs, and if you are requesting PA for a non-preferred drug product, please provide medical necessity for prescribing the non-preferred drug product rather than the preferred drug product.

Prescriber information

Last name* _____ First name* _____ MI _____

NPI* _____ Individual MH Provider ID _____

DEA No. _____ Office Contact Name _____

Address _____ City _____ State _____ Zip _____

E-mail address _____

Telephone No.* _____ Fax No.* _____

* *Required*

Prescribing provider's attestation, signature, and date

I certify under the pains and penalties of perjury that I am the prescribing provider identified in the Prescriber information section of this form. Any attached statement on my letterhead has been reviewed and signed by me. I certify that the medical necessity information (per 130 CMR 450.204) on this form is true, accurate, and complete, to the best of my knowledge. I understand that I may be subject to civil penalties or criminal prosecution for any falsification, omission, or concealment of any material fact contained herein.

Prescribing provider's signature (Signature and date stamps, or the signature of anyone other than the provider, are not acceptable.)

Signature required _____

Printed name of prescribing provider _____ Date _____