Michigan Prior Authorization Request Form
For Prescription Drugs

Instructions

Important: Please read all instructions below before completing FIS 2288.

Section 2212c of Public Act 218 of 1956, MCL 500.2212c, requires the use of a standard prior authorization form when a policy, certificate or contract requires prior authorization for prescription drug benefits.

A standard form, FIS 2288, is being made available to simplify exchanges of information between prescribers and health insurers as part of the process of requesting prescription drug prior authorization. This form will be updated periodically and the form number and most recent revision date are displayed in the top left hand corner.

- Prior authorization requests are defined as requests for pre-approval from an insurer for specified medications or quantities of medications before they are dispensed.
- Prescriber means the term as defined in section 17708 of the Public Health Code, 1978 PA 368, MCL 333.17708.
- Prescription drug means the term as defined in section 17708 of the Public Health Code, 1978 PA 368, MCL 333.17708.
- This form is made available for use by prescribers to initiate a prior authorization request.
- Insurers may request additional information or clarification needed to process a prior authorization request. The prior authorization is not considered granted if the prescriber fails to submit the additional information within 72 hours after the date and time of the original submission of a properly completed prior authorization request.
- In order to designate a prior authorization request for expedited review, a prescriber must certify that applying the 15-day standard review period may seriously jeopardize the life and health of the patient or the patient’s ability to regain maximum function.
Michigan Prior Authorization Request Form for Prescription Drugs  Fax: 858-790-7100

☐ Expedited Review Request: I hereby certify that a standard review period may seriously jeopardize the life or health of the patient or the patient’s ability to regain maximum function.
Physician’s Direct Contact Phone Number ( ) ———— Initials: ————

A) Reason for Request
☐ Initial Authorization Request  ☐ Renewal Request  ☐ DAW

B) Patient Demographics
Is patient hospitalized:  ☐ Yes  ☐ No
Patient Name: ___________________________ DOB: __________________
Patient Health Plan ID: ___________________________
☐ Male  ☐ Female

C) Pharmacy Insurance Plan
☐ Priority  ☐ Magellan  ☐ Blue Cross Blue Shield of Michigan  ☐ HAP  ☐ ________
☐ Total Health Care  ☐ Blue Care Network  ☐ HealthPlus of Michigan  ☐ Meridian Health Plan

D) Prescriber Information
Prescriber Name: ___________________________ NPI: ___________________________ Specialty: ___________________________
DEA (required for controlled substance requests only): ___________________________
Contact Name: ___________________________ Contact Phone: ___________ Contact Fax: ___________
Health Plan Provider ID (if accessible): ___________________________

E) Pharmacy Information (optional)
Pharmacy Name_________________________ Pharmacy Telephone_________________________

F) Requested Prescription Drug Information
Drug Name: ___________________________ Strength: ___________
Dosing Schedule: ___________________________ Duration: ___________________________
Diagnosis (specific) with ICD#: ___________________________
Place of infusion / injection (if applicable): ___________________________
Facility Provider ID / NPI: ___________________________
Has the patient already started the medication? _______ Yes _______ No  If so, when? ___________
G) **Rationale for Prior Authorization** (e.g., information such as history of present illness, past medical history, current medications, etc.; you may also attach chart notes to support your request if you believe they will assist with the review process)

<table>
<thead>
<tr>
<th>Drug Name</th>
<th>Strength</th>
<th>Dosing Schedule</th>
<th>Duration</th>
<th>Adverse Event/Specific Failure</th>
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H) **Failed/Contraindicated Therapies**

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<th>Drug Name</th>
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<th>Dosing Schedule</th>
<th>Duration</th>
<th>Adverse Event/Specific Failure</th>
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I) **Other Pertinent Information** (Optional - to be filled out if other information is necessary such as relevant diagnostic labs, measures of response to treatment, etc.) Please refer to plan's website for additional information that may be necessary for review. Please note that sending this form with insufficient clinical information may result in extended review period or adverse determination.

I represent to the best of my knowledge and belief that the information provided is true, complete and fully disclosed. A person may be committing insurance fraud if false or deceptive information with the intent to defraud is provided.

Physician's Name: ___________________________________________

Physician's Signature: _________________________________________

Date: __________

PA 218 of 1956 as amended requires the use of a standard prior authorization form by prescribers when a patient's health plan requires prior authorization for prescription drug benefits.

*For Health Plan Use Only*

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<th>Request Date:</th>
<th>LOB:</th>
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<tr>
<td>Approved:</td>
<td>Denied:</td>
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<td>Approved By:</td>
<td>Denied By:</td>
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<td>Effective Date:</td>
<td>Reason for Denial:</td>
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<td>Additional Comments:</td>
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Some of the information needed to make a determination for coverage is not specifically requested on the Michigan Prior Authorization Request Form for Prescription Drugs. To avoid delays in reviewing your request, please make sure to include all of the following information.

**Answer questions 1 and 4 for all patients**

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<tr>
<th>Question</th>
<th>Y</th>
<th>N</th>
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<tr>
<td>1. Is the patient age 18 years or older?</td>
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| 2. Does the patient meet one of the following conditions:  
  a. Chronic phase, accelerated phase, or blast phase T3151-positive chronic myeloid leukemia,  
  b. T3151-positive Philadelphia chromosome positive acute lymphoblastic leukemia, or  
  c. Chronic, accelerated, or blast phase Ph+ Acute Lymphocytic Leukemia for whom no other tyrosine kinase inhibitor is indicated? |  |  |
| 3. Will the patient be monitored as clinically indicated for adverse events such as arterial thrombosis, hepatotoxicity, signs and symptoms of congestive heart failure, hypertension, pancreatitis, hemorrhage, fluid retention, cardiac arrhythmias, and myelosuppression? |  |  |
| 4. Will the patient be advised to avoid becoming pregnant while taking Iclusig? | NA |  |