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)

H) Failed/Contraindicated Therapies

<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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<tbody>
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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**

Request Date: ________________________________ LOB: ________________________________

Approved: ________________________________ Denied: ________________________________

Approved By: ________________________________ Denied By: ________________________________

Effective Date: ________________________________ Reason for Denial: ________________________________

Additional Comments: ________________________________
UNIVERSITY OF MICHIGAN – MEDICAL NECESSITY FOR BRAND ANTICONVULSANTS

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.

This form is to allow patients who have been stabilized on a brand anticonvulsant to continue on brand after a generic is introduced. Patients who are initially stabilized on generic anticonvulsants do not need this form.

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<tr>
<td>1.</td>
<td>Is the patient being treated for seizures?</td>
<td>Y</td>
<td>N</td>
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<td>2.</td>
<td>Is this a new medication/therapy start?</td>
<td>Y</td>
<td>N</td>
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<td>3.</td>
<td>Has the patient's seizure been controlled with the brand manufacturer's product?</td>
<td>Y</td>
<td>N</td>
</tr>
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Name of the requested brand (use 1 form per drug):