## 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 by the Department of Insurance and Financial Services 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.

- > This form is made available for use by prescribers to initiate a prior authorization request with the health insurer.
- ➤ 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.
- ➤ Pursuant to MCL 500.2212c, prescribers and insurers must comply with required timeframes pertaining to the processing of a prior authorization request. Insurers may request additional information or clarification needed to process a prior authorization request.
- ➤ The prior authorization is considered granted if the insurer fails to grant the request, deny the request, or require additional information of the prescriber within 72 hours after the date and time of submission of an expedited prior authorization request or within 15 days after the date and time of submission of a standard prior authorization request. If additional information is requested by an insurer, a prior authorization request is considered to have been granted by the insurer if the insurer fails to grant the request, deny the request, or otherwise respond to the request of the prescriber within 72 hours after the date and time of submission of the additional information for an expedited prior authorization request; or within 15 days after the date and time of submission of the additional information for standard prior authorization request.
- The prior authorization is considered void if the prescriber fails to submit the additional information within 5 days after the date and time of the original submission of a properly completed expedited prior authorization request or within 21 days after the date and time of the original submission of a properly completed standard 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.

PRESCRIBERS, PLEASE SUBMIT THIS FORM TO THE PATIENT'S HEALTH PLAN ONLY. Please do not send to the department.

Only provide the physician's direct contact number and initials if you are requesting an Expedited Review Request.

## Michigan Prior Authorization Request Form for Prescription Drugs Fax: 800-424-7648

(PRESCRIBERS SUBMIT THIS FORM TO THE PATIENT'S HEALTH PLAN)

☐ Standard Review Request	
☐ Expedited Review Request: I hereby certify that a standard revies jeopardize the life or health of the patient or the patient's ability to	
Physician's Direct Contact Phone Number:	Initials:
A) Reason for Request  ☐ Initial Authorization Request ☐ Renewal Request ☐ DA	AW
B) Patient Demographics	
Is patient hospitalized: ☐ Yes ☐ No	
Patient Name:	DOB:
Patient Health Plan ID:	
C) Pharmacy Insurance Plan	
☐ Priority ☐ Magellan ☐ Blue Cross Blue Shield of Michigan	☐ HAP ☑ University of Michigan Prescription Drug Plar
☐ Total Health Care ☐ Blue Care Network ☐ HealthPlus of I	Michigan
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: Pharmac	cyTelephone:
F) Requested Prescription Drug Information	
Drug Name: Stre	ength:
Dosing Schedule: Dur	ration:
Diagnosis (specific) with ICD#:	
Place of infusion/injection (if applicable):	
Facility Provider ID/NPI:	
Has the patient already started the medication? ☐ Yes ☐ No	o If so, when?

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relevant diagnost additional information	tic labs, measures ation that may be	of response to treatr necessary for review	ment, etc.) Ple . Please note	ormation is necessary such as ease refer to plan's website for that sending this form with r adverse determination.
Drug Name	Strength	Dosing Schedule	Duration	Adverse Event/Specific Failure
H) Failed/Contrair	ndicated Thera	pies		
		assist with the review		chart notes to support your

G) Rationale for Prior Authorization: (e.g., information such as history of present illness, past



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## University of Michigan – Stivarga® (regorafenib)

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.

Initial Request – Osteosarcoma		
Is the medication being prescribed by an oncologist?	Υ	N
Does the member have a diagnosis of osteosarcoma?	Y	N
Has the member tried an NCCN first line therapy for osteosarcoma?  If yes, please document the specific medications, dates of trials, and clinical outcomes.	Y	N
Initial Request – Rectal/Colon Cancer		
Is the medication being prescribed by an oncologist?	Y	N
Does the member have a diagnosis of advanced or metastatic colon or rectal cancer?	Υ	N
If <b>yes</b> , has the member progressed through all available treatment regimens besides regorafenib or trifluridine/tipiracil?  If <b>yes</b> , please document the specific medications, dates of trials, and clinical outcomes.	Y	N
Initial Request – Glioblastoma		
Is the medication being prescribed by an oncologist?	Υ	N
Does the member have a diagnosis of recurrent glioblastoma?	Υ	N

Continued on the next page.





Member's Last Name:										Member's First Name:													
In	itial R	leque	st – G	IST																			
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da	satinik	(Spry	er exp rcel)? locume															OR			Υ		N
Ini	itial R	leque	st – H	CC																			
Is	the me	edicat	ion bei	ng pr	escrib	ed b	y an o	onco	logist	t?											Υ		N
Is the medication being prescribed by an oncologist?  Does the member have a diagnosis of progressive hepatocellular carcinoma (HCC)?													Υ		N								
Does the member have a diagnosis of progressive hepatocellular carcinoma (HCC)?  The member is <b>NOT</b> a transplant candidate. Is this statement <b>TRUE</b> ?														Υ		N							
			consid												-	does	the r	meml	oer		Y		N
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Ini	itial R	leque	st – S	oft T	issue	Sar	coma	a															
ls t	the me	edicat	ion bei	ng pr	escrib	ed b	y an o	onco	logist	t?											Υ		N
Do	es the	e mem	ber ha	ve a	diagno	osis o	f adv	ance	d/m	etast	atic	pleo	morp	hic r	habd	omyc	sarco	oma?			Υ		N
Do	es the	e mem	ber ha	ve a	diagno	osis o	f a sc	olitar	y fibr	ous t	um	or?									Υ		N
Do	es the	mem	ber ha	ve a	diagno	osis o	f nor	ı-adi <sub>l</sub>	pocyt	ic sa	rco	ma?									Υ		N
Do	es the	mem	ber ha	ve a	diagno	osis o	f ang	iosaı	rcom	a?											Υ		N

Continued on the next page.





Member's Last Name:								Member's First Name:																

Renewal Request		
Does the prescriber attest that the member continues to have a beneficial response to therapy?	Y	N
Is the medication being prescribed by an oncologist?	Υ	N

University of Michigan Prescription Drug Plan Pharmacy Services Portal: <a href="https://umich.magellanrx.com/">https://umich.magellanrx.com/</a>