

Pharmacy Prior Authorization Form

Fax completed form t	o: 877.974.4411 toll free,	or 616.942.8206
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This form applies to:

- Commercial (Traditional)
- Commercial (Individual/Optimized)

Medicaid

This request is:

Urgent (life threatening)
Non-Urgent (standard review)

Urgent means the standard review time may seriously jeopardize the life or health of the patient or the patient's ability to regain maximum function.

Dosing frequency:

Sporanox[®] and itraconazole oral suspension

Member	
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Last Name:		First Name:	
			Gender:
	sician:		
Requesting Provid	ler:	Prov. Phone:	Prov. Fax:
		Contact Name:	
		Date:	
Product Inform	nation		
New request	Continuation request		
Drug product:	Sporanox 10mg/mL oral suspension itraconazole 10mg/mL oral suspension	Start date (or date of next dose): Date of last dose (if applicable):	

Precertification Requirements

Before this drug is covered, the patient must meet all of the following criteria:

- 1. Patient has one of the following conditions:
 - Invasive Aspergillosis
 - Blastomycosis
 - Histoplasmosis
 - Oropharyngeal and esophageal candidiasis
- Inadequate response, intolerable side effect, or contraindication to clotrimazole troches, nystatin suspension, fluconazole <u>and</u> itraconazole oral capsules (oropharyngeal/esophageal candidiasis); or itraconazole oral capsules (all other approved indications).
- 3. Prescribed or recommended by an infectious disease specialist.

Note: Authorization for indications, dosing, or a route of administration not approved by the Food and Drug Administration (FDA) or recognized in CMSaccepted compendia (e.g. DrugDex, AHFS, U.S. Pharmacopeia, and also Clinical Pharmacology for oncology indications only) require supporting evidence for coverage. Please provide two published peer-reviewed literature articles supporting the appropriateness of the drug, the dosing of the drug, or the route of administration to be used for the identified indication.

Priority Health Precertification Documentation

A. What condition is this drug being requested for?

- Invasive Aspergillosis
- Blastomycosis
- Histoplasmosis
- Oropharyngeal and esophageal candidiasis
-] Other the patient's condition is: _____ Rationale for use: _____



B. Was a culture completed?

 Yes 🗌 No

C. Was antifungal susceptibility determined?

Yes (fax results with this request)

No – rationale for use:

D. What antifungals were previously used that were not successful in treating the patient's current infection?

Antifungals used include:

Drug:	Date:	Outcome:
Drug:	Date:	Outcome:
Drug:	Date:	Outcome:

No other antifungals have been used for the patient's current infection.

Note: If adequate blood levels while on itraconazole oral capsules could not be achieved, please provide laboratory value(s) with this request.

Ε. Does the patient have an allergy or contraindication to alternative antifungal therapies?

INO
Yes

163			
Drug:	Date:	Reaction:	
Drug:	Date:	Reaction:	
Drug:	Date:	Reaction:	
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Additional information Notes:

If approved, initial authorization is for a maximum of 3 months (Invasive Aspergillosis, Blastomycosis, Histoplasmosis); oropharyngeal candidiasis (4 weeks); esophageal candidiasis (6 weeks).

Review of precertification requests for indications and/or duration of therapy in the above criteria will be reviewed by a clinical pharmacist and/or medical director.