

Please complete this **entire** form and fax it to: **866-940-7328**. If you have questions, please call **800-310-6826**.
This form contains multiple pages. Please complete all pages to avoid a delay in our decision.
Allow at least 24 hours for review.

Section A – Member Information

First Name:	Last Name:	Member ID:
Address:		
City:	State:	ZIP Code:
Phone:	DOB:	Allergies:
Primary Insurance:	Policy #:	Group #:

Is the requested medication New or Continuation of Therapy? If continuation, list start date: _____
 Is this patient currently hospitalized? Yes No If recently discharged, list discharge date: _____

Section B - Physician Information

First Name:	Last Name:		M.D./D.O.
Address:	City:	State:	ZIP code:
Phone:	Fax:	NPI #:	Specialty:
Office Contact Name / Fax attention to:			

Section C - Medical Information

Medication:	Strength:
Directions for use:	Quantity:
Diagnosis (Please be specific & provide as much information as possible):	ICD-10 CODE:
Is this member pregnant? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, what is this member's due date? _____	

Section D – Previous Medication Trials

Medications	Strength	Directions	Dates of Therapy	Reason for failure / discontinuation

Section E – Additional information and Explanation of why preferred medications would not meet the patient's needs
Please refer to www.uhccommunityplan.com for a list of preferred alternatives

Member First name:	Member Last name:	Member DOB:
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Clinical and Drug Specific Information

- What is the patient’s diagnosis? (check which applies)

- | | |
|---|--|
| <input type="checkbox"/> Aspergillosis
<input type="checkbox"/> Candida Septicemia
<input type="checkbox"/> Candida UTIs
<input type="checkbox"/> Cryptococcus Pulmonary infections
<input type="checkbox"/> Neutropenia Myelodysplastic Syndrome
<input type="checkbox"/> Neutropenic hematologic malignancies
<input type="checkbox"/> Blastomycosis
<input type="checkbox"/> Onychomycosis
<input type="checkbox"/> S. apiospermum
<input type="checkbox"/> Esophageal Candidiasis

<input type="checkbox"/> Other, list diagnosis: _____ | <input type="checkbox"/> Invasive Mucomycosis
<input type="checkbox"/> Candida Endocarditis
<input type="checkbox"/> Cryptococcus Meningitis
<input type="checkbox"/> Oropharyngeal/Esophageal Candidiasis
<input type="checkbox"/> Neutropenic Acute Myeloid Leukemia
<input type="checkbox"/> Graft vs. Host disease
<input type="checkbox"/> Histoplasmosis
<input type="checkbox"/> Candidemia
<input type="checkbox"/> Fusarium spp
<input type="checkbox"/> Immunosuppression secondary to hematopoietic stem cell transplant |
|---|--|

Request for Noxafil Suspension:

- Is the patient’s disease refractory to itraconazole and/or fluconazole?** Yes No
 (If yes, complete Section D above with medication information, including dose, duration, and date of trial)

Request for Onmel:

- Does the patient have a trial and failure or contraindication to terbinafine?** Yes No
 (If yes, complete Section D above with medication information, including dose, duration, date of trial, and reason for discontinuation)

Request for Sporanox Liquid:

- Is there a clinical reason why the oral (capsule) formulation cannot be used?** Yes No
 (If yes, complete Section D above with medication information, including dose, duration, and date of trial)

Request for Sporanox:

- Does the patient have a trial and failure with generic itraconazole?** Yes No

Request for Sporanox/Itraconazole and VFend:

- Is the patient’s onychomycosis due to terbinafine-resistant dermatophytes?** Yes No
 (If yes, complete Section D above with medication information, including dose, duration, and date of trial)
- Is the patient’s diagnosis refractory to fluconazole?** Yes No
 (If yes, complete Section D above with medication information, including dose, duration, and date of trial)

Requests for Non-Preferred Medication: (please refer to www.uhccommunityplan.com)

- Has the patient failed TWO diagnosis-appropriate preferred medications?** Yes No
 (If yes, complete Section D above with medication information, including dose, duration, and date of trial)

Physician Signature: _____ **Date:** _____

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