OPTIMA HEALTH PLAN

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions</u>: <u>The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request</u>. All other information may be filled in by office staff; fax to <u>1-800-750-9692</u>. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. <u>If information provided is not complete, correct, or legible, authorization may be delayed.</u>

<u>Drug Requested:</u> Brexafemme[®] (ibrexafungerp)

| ME | MBER & PRESCRIBER I | NFORMATION: Authorization may be delayed if incomplete. |
|------------------|---------------------------------|--|
| Meml | ber Name: | |
| Member Optima #: | | |
| Presc | riber Name: | |
| | | Date: |
| Office | e Contact Name: | |
| Phone Number: | | Fax Number: |
| DEA | OR NPI #: | |
| | | norization may be delayed if incomplete. |
| Drug | Form/Strength: | |
| Dosing Schedule: | | |
| | | ICD Code: |
| Weight: | | Date: |
| supp | | below all that apply. All criteria must be met for approval. To ntation, including lab results, diagnostics, and/or chart notes, must be |
| Diag | gnosis: Vulvovaginal Cand | lidiasis (VVC), acute infection |
| Reco | ommended Dosing: 300 mg ever | y 12 hours for 1 day (2 doses) |
| Len | gth of Authorization: Date | of Service, one-time fill |
| | Member is post-menarchal | |
| | Provider has confirmed that the | member is not pregnant |
| | • | for acute, uncomplicated vulvovaginal candidiasis (VVC) (please ation or medical chart notes to confirm diagnosis i.e., urinalysis, 10% KOH) |
| | | zole at the recommended dosage of 150 mg as a single dose for the claims history and chart notes must confirm failure) |

(Continued on next page)

| | ☐ Trial and failure of two topical agents (suppository inserts/ovules/creams) for the treatment of VVC (pharmacy claims history and chart notes must confirm failure) | | |
|-----------------------------------|---|---|--|
| | | Gynazole-1 vaginal cream 2 % | |
| | | Terconazole vaginal cream 0.4 %, 0.8 % | |
| | | Terconazole vaginal suppository 80 mg | |
| | | OTC products: tioconazole ointment 6.5%, miconazole suppository 100 mg/200 mg, clotrimazole cream 1%, 2% /100 mg suppository | |
| Diag | gno | sis: Recurring Vulvovaginal Candidiasis (RVVC) | |
| Reco | mm | nended Dosing: 300 mg every 12 hours for 1 (2 doses); repeat monthly for a total of 6 months | |
| Length of Authorization: 6 months | | | |
| | Me | ember is post-menarchal | |
| | Pro | ovider has confirmed that the member is not pregnant | |
| | Member is currently experiencing signs and symptoms consistent with an acute episode of VVC (e.g., vulvovaginal pain, pruritis or irritation, abnormal vaginal discharge), AND it is a laboratory confirmed VVC episode (please include laboratory documentation or medical chart notes to confirm diagnosi i.e., urinalysis, microscopic examination via 10% KOH, culture) | | |
| | Member has a history of recurring VVC (RVVC) (please include past medical history notes recording RVVC, defined as \geq 3 episodes of vulvovaginal candidiasis (VVC) in a 12-month period) | | |
| | dos no | ember remains symptomatic and culture positive after therapy with fluconazole, completing a 6-month sing regimen as follows unless intolerant or contraindicated (please include medical chart/progress tes and laboratory results; pharmacy claims history and chart notes must confirm failure, | |
| | int | colerance or contraindication to therapy): | |
| | | 100, 150 or 200 mg oral dose of fluconazole every third day for a total of 3 doses (days 1, 4 and 7) | |
| | | Followed by oral fluconazole (100, 150 or 200 mg oral dose) weekly for 6 months as the maintenance regimen | |
| | | | |

Not all drugs may be covered under every Plan

If a drug is non-formulary on a Plan, documentation of medical necessity will be required.

**Use of samples to initiate therapy does not meet step edit/ preauthorization criteria. **

*Previous therapies will be verified through pharmacy paid claims or submitted chart notes. *

REVISED/UPDATED: 10/29/2021; 11/1/2021; 3/9/2023

^{*}Approved by Pharmacy and Therapeutics Committee: 9/16/2021; 2/16/2023